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                 feature for sorting BLAST answer sets
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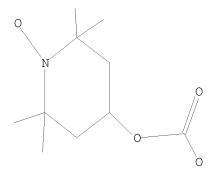
chain nodes : 7 14 ring nodes : 1 2 3 4 5 6 ring/chain nodes : 8 9 10 11 12 15 16 chain bonds : 2-9 2-10 3-8 4-11 4-12 6-7 7-14 14-15 14-16 ring bonds : 1-2 1-6 2-3 3-4 4-5 5-6 exact/norm bonds : 1-2 1-6 2-3 3-8 5-6 6-7 7-14 14-15 14-16 exact bonds : 2-9 2-10 3-4 4-5 4-11 4-12 isolated ring systems : containing 1 :

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# L1 STRUCTURE UPLOADED

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=> s 11

SAMPLE SEARCH INITIATED 15:37:54 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 5 TO ITERATE

100.0% PROCESSED 5 ITERATIONS 1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*

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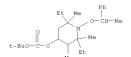
PROJECTED ITERATIONS: 5 TO 234

PROJECTED ANSWERS: 1 TO 80

L2 1 SEA SSS SAM L1

=> d scan

L2 1 ANSWERS REGISTRY COPYRIGHT 2009 ACS on STN
IN Carbonic acid, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4piperidinyl 1,1-dimethylethyl ester
NF C25 H41 N O4



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=> 1

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100.0% PROCESSED 93 ITERATIONS

33 ANSWERS

SEARCH TIME: 00.00.01

L3 33 SEA SSS FUL L1

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L1 STRUCTURE UPLOADED

L2 1 S L1

L3 33 S L1 FULL

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=> s 13

L4 67 L3

=> d cbib abs hitstr 1-YOU HAVE REQUESTED DATA FROM 67 ANSWERS - CONTINUE? Y/(N):y

L4 ANSWER 1 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:1315916 Document No. 151:5500670 Tetraarylphosphonium Salts as Soluble Supports for Oxidative Catalysts and Reagents. Roy, Marie-Noelle; Poupon,

on, Jean-Christophe; Charette, Andre B. (Departement de Chimie, Universite de Montreal, Montreal, QC, H3C 3J7, Can.). Journal of Organic Chemistry, 74(22), 8510-8515 (English) 2009. CODEN: JOCEAH. ISSN: 0022-3263. OTHER

R
SOURCES: CASREACT 151:550067. Publisher: American Chemical Society.
Tetraarylphosphonium (TAP)-supported DMSO, TEMPO, and
(diacetoxy)iodobenzene reagents were synthesized and used for the ovidation

of alcs., including Swern oxidation, and for the  $\alpha$ -acetoxylation of ketones. By taking advantage of the predictable solubility properties of the

TAP unit, simple precipitation and filtration of the phosphonium moiety

permitted

complete separation of the desired products. It was demonstrated that

these
reagents could be recycled directly when used in catalytic processes and
following regeneration when used in stoichiometric processes.

IT 867023-62-9P
RL: CAT (Catalyst use); RCT (Reactant); SPN (Synthetic preparation); PREP
(Preparation); RACT (Reactant or reagent); USES (Uses)
(preparation of recyclable tetraarylphosphonium-supported DMSO,
TEMPO, and
iodobenzene and use as reagent/catalysts in the oxidation of alcs. and
acetoxylation of ketones)
RN 867023-62-9 CAPLUS
C1-Pieridinyloxy, 2,2,6,6-tetramethyl-4-[[[4(triphenylphosphonio)phenyl]methoxy]carbonyl]oxy]-, perchlorate (1:1)

(CA

INDEX NAME)

CM 1

CRN 867023-61-8 CMF C35 H38 N O4 P

CM 2

CRN 14797-73-0 CMF Cl O4

14 ANSWER 2 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2009:1139409 Document No. 151:359933 Synthetic resin composition and automotive interior/exterior material comprising the same. Mizokawa, Shigeo; Negishi, Yoshinori (Adeka Corporation, Japan). PCT Int. Appl. v
2009:113389 A1 2009:0917, 32pp. DESIGNATED STATES: W: AE, AG, AL, AM, AA, AT, AJ, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, II, IN, IS, JP, KE, KG, RM, KN, KP, KR, KZ, LA, LC, LK, LE, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MM, MX, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, PH, FL, FT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NI, NO, PT, SE, SN, TD, TG, TR. (Japanese). CODEN: PIXXD2. APPLICATION: WO
2009—JF53341
20090225. PRIORITY: JP 2008-59576 20080310.

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

A synthetic resin composition with good weather resistance useful for automotive interior/exterior material comprising 100 parts synthetic resin, 0.01-20 parts component (A) and 0.01-20 parts component (B), wherein the component (A) is a hindered amine compound represented by general formula I (R = 1-30 alkyl, C2-30 alkenyl; n = 1-6; R1 = C1-22 alkyl, C2-22 alkyl, C2-22 alkyl, C2-22 alkyl, R2 = (C1-30 alkyl, R) = Too book (C1-21 linear or branched alkyl; n = 2-6; X = -C(:O)-), and the component (B) is a sulfur-containing antioxidant. 705257-84-7P

705257-84-7P
RL: IMF (Industrial manufacture); MOA (Modifier or additive use); PREP (Preparation); USES (Uses)
(synthetic resin composition for automotive interior/exterior

material)

:141)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA
INDEX NAME)

L4 ANSWER 1 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

IT 867023-64-1P RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of recyclable tetraarylphosphonium-supported DMSO, TEMPO, and

o, and iodobenzene and use as reagent/catalysts in the oxidation of alcs. and acetoxylation of ketones) 867023-64-1 CAPLUS Phosphonium, [4-[[[[(1-hydroxy-2,2,6,6-tetramethyl-4-piperidinyl)oxy]carbonyl]oxy]methyl]phenyl]triphenyl-, perchlorate (1:1) (CA INDEX NAME)

CM 1

CRN 867023-63-0 CMF C35 H39 N O4 P

CM 2

CRN 14797-73-0 CMF C1 04

L4 ANSWER 3 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:1131615 Document No. 151:314742 Woody synthetic resin composition having improved weather resistance and molded body thereof. Fukushima Mitsuru; Mizokawa, Shigeo; Mizu, Masumi (Adeka Corp., Japan). Jpn. Ko Tokkyo Koho JP 2009209205 A 20090917, 15pp.; Chemical Indexing Equivalent to 151:314717 (WO) (Japanese). CODEN: JKXXAF. APPLICATION: JP 2008-51060 20080229.

Disclosed are a woody synthetic resin composition having excellent

resistance, and a molded body of the woody synthetic resin compn (e.g., ethylene-propylene copolymer). The woody synthetic resin composition is obtained by blending 5-200 parts by mass of wood flour and 0.01-5 parts

mass of a hindered amine compound having a partial structure represented

the general formula I per 100 parts by mass of a synthetic resin (R1, R2, R3 and R4 independently represent an alkyl group having 1-4 carbon atoms; and R represents an alkyl group having 1-18 carbon atoms, an alkyl group substituted by a hydroxy group or a cycloalkyl group having 5-8 carbon atoms). atoms). 705257-84-7

RL: MOA (Modifier or additive use); USES (Uses) (woody synthetic resin composition having improved weather resistance

and molded body thereof) 705257-84-7 CAPLUS

ANSWER 4 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1131570 Document No. 151:382449 Polylactic acid resin composition
containing hindered amine having excellent weather resistance. Negishi,
Yoshinori; Mizokawa, Shigeo (Adeka Corp., Japan). Jpn. Kokai Tokkyo Koho
JP 2009209177 A 20090917, 15pp.; Chemical Indexing Equivalent to
151:314707 (NO) (Japanese). CODEN: JKXXAF. APPLICATION: JP 2008-50630
20080229.

131:314/W/ (WO) (Japanese). CODEN: JKXXAF. APPLICATION: JP 2008-50630 20080229.

AB The invention relates to a polylactic acid resin composition obtained by blending 0.005-30 parts of a hindered amine compound, which has a partial structure represented by formula -C(R1) (R2)-R(08)-C(R3) (R4)-, per 100 parts of a polylactic acid resin; wherein R1, R2, R3 and R4 independently represent an alkyl group having 1-18 carbon atoms; and R represents an alkyl group having 1-18 carbon atoms which may be substituted by a hydroxyl group, an acyl group having 1-18 carbon atoms or a cycloalkyl group having 5-8 carbon atoms. Thus, a polylactic acid resin composition suppressing crystallization and having excellent weather resistance was prepared by mix-dissolving polylactic acid resin (Lacea H 100) 1.25 g and a hindered amine 0.375 mg in methylene chloride, and by casting the solution to give a cast film having a crack conserving the second conserving the solution to

L4 ANSWER 5 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

ANSWER 5 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1081125 Document No. 151:314717 Woody synthetic resin composition having improved weather resistance and molded body thereof. Fukushima, Mitsuru; Mizokawa, Shiqeo; Mizu, Masumi (Adeka Corporation, Japan). PCT Int. Appl. Wo 2009107502 Al 2009903, 25pp.; Chemical Indexing Equivalent to 151:314742 (JP) DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU,

BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, IN, IS, KE, KG, EM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, MD, ME, MG, MK, MN, MM, MX, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, FT, BO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TT; RN: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NL, NO, FT, SE, SN, TR. (Japanese). CODEN: PIXND2. APPLICATION: WO 2009-JP52539 2 PRIORITY: JP 2008-51060 20080229.

Disclosed are a woody synthetic resin composition having excellent

rer resistance, and a molded body of the woody synthetic resin compn (e.g., ethylene-propylene copolymer). The woody synthetic resin composition iobtained by blending 5-200 parts by mass of wood flour and 0.01-5 parts

mass of a hindered amine compound having a partial structure represented

the general formula I per 100 parts by mass of a synthetic resin (R1, R2, R3 and R4 independently represent an alkyl group having 1-4 carbon atoms; and R represents an alkyl group having 1-18 carbon atoms, an alkyl group substituted by a hydroxy group or a cycloalkyl group having 5-6 carbon

IT

RL: MOA (Modifier or additive use); USES (Uses)
(woody synthetic resin composition having improved weather resistance and

molded body thereof) 705257-84-7 CAPLUS

/UD/20/-84-/ CAPLUS 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 6 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

2009:1080458 Document No. 151:314707 Polylactic acid resin composition containing hindered amine having excellent weather resistance. Negishi, Yoshinori; Mizokawa, Shigeo (Adeka Corp., Japan). PCT Int. Appl. WO
2009107504 Al 20099093, 25pp.; Chemical Indexing Equivalent to 151:382449
(JP) DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BB, BR, BW, BY, BY, BZ, CA, CR, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, KE, KG, FM, KN, KP, KR, KZ, LA, LC, LK, LK, LS, LT, LU, LY, MA, DM, DM, MG, MK, MY, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, PB, PL, FT, RO, RS, RU, SC, SD, SE, SG, SK, SI, SM, ST, SY, SY, TJ, TM, TN, FT, TF; RN AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FT, FR, GA, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NI, NO, PT, SE, SN, TD, TG, TR.
(Japanese). CODEN: PIXXD2. APPLICATION: WO 2009-JP52563 20090216.
PRIORITY: JP 2008-50630 20080229

AB The invention relates to a polylactic acid resin composition obtained by blending 0.005-30 parts of a hindered amine compound, which has a partial structure represented by formula — C(R1) (R2)—N(OR)—C(R3) (R4)—, per 100 parts of a polylactic acid resin; wherein R1, R2, R3 and R4 independently represent an alkyl group having 1—18 carbon atoms which may be substituted by a hydroxyl group having 1—18 carbon atoms which may be substituted by a hydroxyl group having 1—18 carbon atoms which may be substituted by a hydroxyl group having 1—18 carbon atoms which may be substituted by a mydroxyl group an acyl group having 1—18 carbon atoms or a cycloalkyl group having 5–8 carbon atoms. Thus, a polylactic acid resin composition suppressing crystallization and having excellent weather resistance was prepared by mix—dissolving polylactic acid resin (Lace H 100) 1.25 g and a hindered

prepared by mix-dissolving polylactic acid resin (Lacea H 100) 1.25 g and a hindered amine 0.375 mg in methylene chloride, and by casting the solution to

a cast film having a crack generation time of 1440 h and a hazing rate caused by crystallization of 6.6% after 360 h. 705257-84-7

TT 705257-84-7

RL: MOA (Modifier or additive use); USES (Uses)
(production of polylactic acid resin composition containing hindered amine having
excellent weather resistance)
RN 705257-84-7 CAPLUS
CN 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSMER 7 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:1077233 Document No. 151:315867 Sealing compositions containing hydrolyzable silyl-containing polymers with suppressed peeling from

hydrolyzable silyl-containing polymers with suppressed peeling from glass
Oka, Toruy Murase, Masaaki, Nakayama, Yoshimitsu (Sunstar Engineering Inc., Japan). Jpn. Kokai Tokkyo Koho JP 2009197177 A 20090903, 16pp. (Japanese). CODEN: JRXXAF. APPLICATION: JP 2008-42699 20080225.
AB The compns., useful for sealing light-transmitting parts (e.g., glass), preferably with silicone sealants, contain the polymers and hindered anines bearing N-Cl-20-alkoxy)-2,2,6,6-tetramethyl-4-piperidyl groups. Thus, applying a sealant containing Epion EP 5058 (alkoxysilyl-terminated polyisobutylene) 160, Epikote 828 (epoxy resin) 6, Tinuvin 123 (HALS) 1, and S 1000 (silyl-crosslinkable reactive plasticizer) 40 parts to a primer-coated Al alloy sheet and bonding it to a float glass sheet via Penguinseal 2520 (silicone sealant) gave a test piece showing good interlayer adhesion and weather resistance.

17 705257-84-7 863984-48-9 1185256-91-0
1185256-92-1
RL: MOA (Modifier or additive use); USES (Uses)
(light stabilizer; sealing compns. containing hydrolyzable silyl-containing polymers and N-alkoxy HALS with suppressed peeling from glass)
NN 705257-84-7 CAPLUS
CN 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

863984-48-9 CAPLUS
4-Fiperidinol, 1-(heptadecyloxy)-2,2,6,6-tetramethyl-, carbonate (2:1)
(ester) (9C1) (CA INDEX NAME)

1185256-91-0 CAPLUS
4-Piperidinol, 1-butoxy-2,2,6,6-tetramethyl-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 8 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:916510 Document No. 151:222190 UV absorber compositions, light-resistant synthetic polymer compositions containing them, and films, sheets, and coatings comprising them. Yoshitake, Toshitaka; Tanaka, Tomoki; Kamimoto, Tetsuo (Adeka Corp., Japan). Jpn. Kokai Tokkyo Koho JI 2009167416 A 20090730, 25pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2009-35773 20090218.

The UV absorber compns. contain triazine compds. of I (R1 = C1-12 alkyl, C3-8 cycloalkyl, C3-8 alkenyl, C6-18 aryl, C7-18 alkylaryl, C7-18 arylalkyl; R2 = H, C1-6 alkyl, C3-8 alkenyl; R3 = H, OH; R4 = H, OR1) and hindered amines of II (R5 = C1-12 alkyl, C1-12 alkoxy; R6 = carbonate, mono-, di-, tri-, or tetravalent organic carboxylic acid residue; n =

Thus, a composition comprising triacetyl cellulose (LT 35) 100, I (RI = CH2CH2CH2CO2Me, R2 = Me, R3 = OH, R4 = OR1) 0.2, and II (R5 = OC11H23, R6 = carbonate, n = 2) 0.2 part was east to give a film showing change in yellowness index (AVI) 0.22, 0.49, and 0.78 after a weathering test (83°, no rain, carbon arc) for 120, 240, and 360 h, resp. 705257-84-7
RI: MOA (Modifier or additive use); USES (Uses)
(UV absorber; UV absorber compns. for light-resistant synthetic mer

ner films, sheets, and coatings) 705257-84-7 CAPLUS 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 7 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

1185256-92-1 CAPLUS
Carbonic acid, C,C'-1,2-ethanediyl
C,C'-bis[2,2,6,6-tetramethyl-1-(undecyloxy)-4-piperidinyl] ester (CA
RNDEX NAME)

PAGE 1-B

- (CH2)10-Me

L4 ANSWER 8 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

ANSWER 9 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :795991 Document No. 151:79136 UV stabilized crosslinkable polyolefin compositions comprising acidic silanol condensation catalysts. Nylander, Perry (Borealis Technology Cy, Finland). PCT Int. Appl. WO 2009080235 A1 20090702, 33pp.; Chemical Indexing Equivalent to 151:79085 (EP) DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH,

DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BR, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FIT, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MN, MX, MZ, NA, NG, NI, NO, NZ, CM, FG, PH, FL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SY, SY, TJ, TM, TN, TR, KN: AT, BE, BF, BJ, CT, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GK, IE, IS, IT, LU, MC, ML, MK, MT, NE, NL, NO, PT, SE, SN, TD, TG, TR. (Emglish). CODEN: PIXXD2. APPLICATION: WO 2008-EP10653 20081215. PRIORITY: EP 2007-24836 20071220.

AB A polyolefin composition comprises a crosslinkable polyolefin with hydrolyzable silane groups, and a silanol condensation catalyst, wherein the composition is characterized by excellent curing properties, namely, the maximum torque AF is > 40 Nm and the crosslinking speed is > 0.1 Nm/s in the ice test, and retention of > 60% of the elongation at break after 500 h in SEPAP UV exposure. Preferably, the polyolefin composition further comprises at least one UV stabilizer that is acidic (pH ≤ 6.2, measured at 20-25° in 1% suspension). The polyolefin composition can be used for production of tubes and insulating layers for elec. wires and cables.

IT 705257-84-7

RL: MOA (Modifier or additive use); USES (Uses)

705257-84-7
RL: MOA (Modifier or additive use); USES (Uses)
(ADK-Stab LA 81; UV stabilized crosslinkable polyolefin compns. comprising acidic silanol condensation catalysts)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 11 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009;671515 Document No. 150:565321 Resin composition and resin molded article with good weather resistance. Miscokawa, Shigeo; Negishi, Yoshinori (Adeka Corp., Japan). Jpn. Kokai Tokkyo Koho JP 2009120723 A 20090604, 14pp.; Chemical Indexing Equivalent to 150:565158 (MO) (Japanese). CODEN: JKXXAF. APPLICATION: JP 2007-296312 20071115.

Disclosed is a resin composition containing zinc sulfide and having

Hent weather resistance. Also disclosed is a resin molded article obtained by using such a resin composition Specifically disclosed is a resin

obtained by blending 0.1-20 parts by weight of zinc sulfide (A) and 0.01-20  $\,$ 

parts by weight of a hindered amine compound (B) having a structure

parts by weight of a hindered amine compound (B) having a structure represented by the general formula I (R independently representing an alkyl group having 1-18 carbon atoms which may be substituted by a hydroxyl group, an acyl group having 1-18 carbon atoms, or a cycloalkyl group having 5-8 carbon atoms), per 100 parts by weight of a resin.

IT 705257-84-7

RL: MOA (Modifier or additive use); USES (Uses) (resin composition and resin molded article with good weather resistance) RN 705257-84-7 CAPLUS

4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

ANSWER 10 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :756280 Document No. 151:79085 UV stabilized crosslinkable polyolefin compositions comprising acidic silanol condensation catalysts. Nylander, Perry (Borealis Technology Gy, Finland). Eur. Pat. Appl. EP 2072568 A1 20090624, 17pp.; Chemical Indexing Equivalent to 151:79136 (WO) DESIGNATED STATES: R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR,

GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, FL, FT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS. (English). CODEN: EEXXDW. APPLICATION: EP 2007-24836 20071220.

A polyolefin composition comprises a crosslinkable polyolefin with olyrable

olyzable silane groups, and a silanol condensation catalyst, wherein the osition is characterized by excellent curing properties, namely, the maximum torque  $\Delta F$  is > 40 km and the crosslinking speed is > 0.1 km/s in the ice test, and retention of > 60% of the elongation at break after 500 h in SEPAP UV exposure. Preferably, the polyolefin composition further rises at

SEPAP UV exposure. Preferably, the polyolefin composition further comprises at least one UV stabilizer that is acidic (pH ≤ 6.2, measured at 20-25° in 1% suspension). The polyolefin composition can be used for production of tubes and insulating layers for elec. wires and cables.

IT 705257-84-7, ADK Stab LA 81 RL MOA (Modifier or additive use); USES (Uses)

(ADK Stab LA 81; UV stabilized crosslinkable polyolefin compns. comprising acidic silanol condensation catalysts)

RN 705257-84-7 CAPLUS

CN 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 12 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:655567 Document No. 151:57243 Compound polymerization inhibitor and

application. Zhang, Tianlin; Liu, Lin; Zhou, Jiexing (Huaihai Institute of Technology, Peop. Rep. China). Faming Zhuanli Shenqing Gongkai Shuomingshu CN 101440286 A 20090527, 6pp. (Chinese). CODEN: CNXXEV. APPLICATION: CN 2010-176117 20081103.
The title polymerization inhibitor comprises components A, B and C at a

mass ratio of (10-50):(20-80):(30-70). Component A comprises tris(1-oxy-2,2,6,6-tetramethylpiperidine-4-oxy)phosphorite (TTMPP), bis(1-oxy-2,2,6,6-tetramethylpiperidine-4-oxy)carbonate (BTMBFC) and (1-oxy-2,2,6,6-tetramethylpiperidine-4-oxy)carbonate (BTMBFC) and comprises 4,6-dinitro-2-sec-Bu phenol (DNBF), 4,6-dinitro-p-cresol (DNFC) and 4,6-dinitro-p-cresol (DNFC). Component C comprises 3,5-dimethylthio-2,4-diamino toluene, 2,6-dimethylthio-1,4-diaminobenzene and 4,6-dimethylthio-1,4-diaminobenzene and 1,6-dimethylthio-1,3-diaminobenzene. The invented polymerization inhibitor is

inhibitor is

liquid at low temperature, and used in production or purification of

one, divinylbenzene, p-chloromethyl styrene, 2-vinyl pyridine, 4-vinyl pyridine, polyol acrylate, etc. 6146-58-3

6146-58-3
RL: CAT (Catalyst use); USES (Uses) (compound polymerization inhibitors) 6146-58-3 CAPLUS

win-po-3 CAPLUS 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 13 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2009:618497 Document No. 150:5651580 Resin composition and resin molded
article with good weather resistance. Mizokawa, Shigeo; Negishi,
Yoshinori (Adeka Corporation, Japan). PCT Int. Appl. WO 2009063708 A1
20090522, 24pp.; Chemical Indexing Equivalent to 150:565321 (JP)
DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH,

BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, KE, KG, RM, RN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MM, MX, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, PH, PL, PT, KO, RS, KU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT; RM: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, LE, IS, IT, LU, MC, ML, MR, MT, NE, NL, NO, PT, SE, SN, TD, TG, TR. (Japanese). CODEN: PIXXID: APPLICATION: WO 2008—JP68293 20081008. PRIORITY: JP 2007-296312 20071115.

AB Disclosed is a resin composition containing zinc sulfide and having excellent weather resistance. Also disclosed is a resin molded article obtained by using such a resin composition Specifically disclosed is a resin

composition obtained by blending 0.1-20 parts by weight of zinc sulfide (A) and 0.01-20

0.01-20
parts by weight of a hindered amine compound (B) having a structure represented
by the general formula I (R independently representing an alkyl group having 1-18 carbon atoms which may be substituted by a hydroxyl group, an acyl group having 1-18 carbon atoms, or a cycloalkyl group having 5-8 carbon atoms), per 100 parts by weight of a resin.

TT 705257-84-7

/US20/-64-/ RL: MOA (Modifier or additive use); USES (Uses) (resin composition and resin molded article with good weather

resistance
RN 705257-84-7 CAPLUS
CN 4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 14 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009;583530 Document No. 150;540727 Synthetic resin composition useful for automotive interior/exterior material. Misokawa, Shigee; Negishi, Yoshinori (Adeka Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2009102556 A 20090514, 18pp.; Chemical Indexing Equivalent to 150;473711 (MO) (Japanese). CODEN: JKXXAF. APPLICATION: JP 2007-277139 20071025.

II

The composition, having good weather resistance, contains 100 parts

AB The composition, having good weather resistance, contains 100 parts synthetic resin, 0.01-20 parts a hindered amine compound (A) having a partial structure represented by a general formula I, and 0.01-20 parts a benzoate compound (B) represented by a general formula II, wherein R1-R4 = C1-4 alkyl, R = optionally hydroxy-substituted C1-18 alkyl, or C5-8 cycloalkyl;

R5, R6 = C1-8 alkyl, R7 = C1-30 alkyl, C6-30 aryl, C7-30 alkyl, C7-30 dialkyl aryl, C7-30 trialkyl aryl, or C7-30 aryl alkyl; and the ratio of A/B is 1:1-1:5. Thus, ethylene-propylene copolymer 85, talc 15, gray pigment 3.0, pentaerythritol 3-(4-hydroxy-3,5-di-tert-butylphenyl) propionate 0.1, tris(2,4-di-tert-butylphenyl) phosphite 0.1, calcium stearate 0.1, l-undecyloxy-2,2,6,6-tetramethyl-4-piperidinol carbonate 0.1, and hexadecyl 3,5-di-tert-butyl-4-hydroxybenzoate 0.1

s
were kneaded at 230° to give a title composition
705257-84-7
RL: NOA (Modifier or additive use); USES (Uses)
(synthetic resin composition useful for automotive interior/exterior
material)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA

L4 ANSWER 13 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

ANSWER 14 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN INDEX NAME)

(Continued)

Me-(CH2)10-0 O- (CH2)10-Me

L4 ANSWER 15 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2009:519143 Document No. 150:4737110 Synthetic resin composition useful for automotive interior/exterior material. Mirokawa, Shigeo; Negishi, Yoshinori (Adeka Corporation, Japan). PCT Int. Appl. WO 2009054267 Al 20090430, 26pp.; Chemical Indexing Equivalent to 150:540727 (JP) DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, RB

BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, KE, KG, RM, RN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MM, MX, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, PH, PL, PT, KO, RS, KU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT; RM: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, LE, IS, IT, LU, MC, ML, MR, MT, NE, NL, NO, PT, SE, SN, TD, TG, TR. (Japanese). CODEN: PIXXID: APPLICATION: WO 2008—JP68289 20081008. PRIORITY: JP 2007-277139 20071025.

AB The composition, having good weather resistance, contains 100 parts synthetic

metric resin, 0.01-20 parts a hindered amine compound (A) having a partial structure represented by a general formula I, and 0.01-20 parts a

pate compound (B) represented by a general formula II, wherein R1-R4 = C1-4 alkyl, R = optionally hydroxy-substituted C1-8 alkyl, or C5-8 cycloalkyl; R5, R6 = C1-8 alkyl, R7 = C1-30 alkyl, C6-30 aryl, C7-30 alkyl, C7-30 dialkyl aryl, c7-30 trialkyl aryl, or C7-30 aryl alkyl; and the ratio of A/B is 1:1-1:5. Thus, ethylene-propylene copolymer 85, talc 15, gray pigment 3.0, pentacythritol 3-(4-hydroxy-3,5-di-tert-butylphenyl)propionate 0.1, tris(2,4-di-tert-butylphenyl) phosphite 0.1,

L4 ANSWER 16 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2009:426827 Document No. 150:4271930 Dye-sensitized solar cell containing nitroxy radical compound in electrolyte layer. Nanbu, Yoko (Adeka Co., Ltd., Japan). Jpn. Rokai Tokkyo Koho JP 2009076369 A 20090409, 14pp. (Japanese). COEDN: JKXXAF. APPLICATION: JP 2007-245519 20070921.

AB In the dye-sensitized solar cell having an electrode substrate, a transparent conductive layer, a dye-adsorbed metal oxide layer, an electrolyte layer, another transparent conductive layer, and a counterelectrode in this order, the electrolyte layer contains nitroxy radical compound (acting as redox mediator). The solar cell has improved electromotive force, maximum output, and cycle performance.

IT 28949-43-0

EL: CAT (Catalyst use); USES (Uses) (redox mediators; dye-sensitized solar cell containing nitroxy radical compound in electrolyte layer)

EN 28949-43-0 CAPUS

CN 1-Piperidinyloxy, 4-[(ethoxycarbonyl)oxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

ANSWER 15 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued) calcium stearate 0.1, 1-undecyloxy-2,2,6,6-tetramethyl-4-piperidinol carbonate 0.1, and hexadecyl 3,5-di-tert-butyl-4-hydroxybenzoate 0.1

were kneaded at 230° to give a title compn. 705257-84-7

705257-84-7

RL: MCA (Modifier or additive use); USES (Uses)
(synthetic resin composition useful for automotive interior/exterior material)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 17 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2008:1179870 Document No. 149:4033500 Vinyl chloride polymer waterproof sheet with good weather resistance. Mitsudera, Taro; Sengoku, Tadashi; Yonezawa, Yutaka (Adeka Corporation, Japan) PCTI Int. Appl. WO
2008117575
A1 20081002, 20 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CR, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, CM, GT, HN, HR, HU, IL, ILI, III, IS, KE, KG, KM, KN, KP, KE, KZ, LA, LC, LK, LE, LET, LU, LY, MA, MD, ME, MG, MK, MN, MW, MY, MZ, NA, NG, NI, NO, NZ, CM, PG, PL, PT, FO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TZ, TZ, TW. AT, BE, BF, BT, CT, CG, CH, CI, CM, CY, DE, DK, ES, FI, TG, GR, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NL, NO, PT, SE, SN, TD, TG, TR. (Japanese). COORNIPIXMEZ. APPLICATION: WO 2008-JP52240 20080212. PRIORITY: JP 2007-80780 20070327.

The waterproof sheet is made of a vinyl chloride resin composition

AB The waterproof sheet is made of a vinyl chloride resin composition comprising

(A) 100 parts a vinyl chloride resin, (B) 5-100 parts a plasticizer, (C) 0.05-5 parts a hindered amine compound I (Rl, R2 = C4-20 alkyl, C5-6 cycloalkyl), and (D) 0.05-5 parts a benzotriazole UV absorber II (R3 = C1-12 alkyl).

IT 705257-84-7

R1: MOA (Modifier or additive use); USES (Uses)

(vinyl chloride polymer waterproof sheet with good weather resistance)
RN 705257-84-7 CAPLUS

4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 17 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

L4 ANSWER 19 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2008:801631 Document No. 149:1303690 Multifilament, monofilament, nonwoven or tape containing hindered amines as light stabilizers. Mueller,
Daniel;

Pauquet, Jean-Roch; Judge, Anthony; Meyer, Hanspeter (Ciba Holding Inc., Switz.). PCT Int. Appl. No 2008077830 A2 20080703, 55pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CH, CO, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, CE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, EM, KN, KP, KR, KZ, LA, LC, LK, LE, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MN, MY, MY, MN, NA, NN, NN, CZ, CM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, FW: AT, BE, BF, BJ, CF, GG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NI, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: NO 2007-EP64023 20071217. PRIORITY: EP 2006-127222

A multifilament, a monofilament, a non-woven or a tape, each having 1 - 2000 Denier per filament and a draw ratio of 1:2 - 1:11 and each made of AB

composition containing the components (A) a polyolefin, (B) 2 hindered

composition containing the components (A) a polyolefin, (B) 2 hindered amines,
e.g., I and II (b1 = 2-20), and optionally (C) one or more inorg. and/or organic pigments.

IT 705257-84-7
RL: MOA (Modifier or additive use); USES (Uses)
(light stabilizer; manufacture of multifilament, monofilament, nonwoven or
tape containing hindered amine additives)

RN 705257-84-7 CAPLUS
A-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 18 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2008:1069425 Document No. 149:3332270 Hindered alkoxyamine-containing acidic

ic
polymer compositions with long-lasting weather resistance. Negishi,
Yoshinori; Dobugawa, Shigeo (Adeka Co., Ltd., Japan). Jpn. Kokai Tokkyo
Koho JP 2008202005 A 20080904, llpp. (Japanese). CODEN: JKXXAF.
APPLICATION: JP 2007-42883 20070222.

AB Title compns., useful for agricultural films, etc., contain 100 parts acid

acid group-containing polymers and 0.005-30 parts hindered amines having partial

partial structures CH2CR1R2N(OR)CR3R4CH2 [R1-4 = C1-4 lower alkyl; R = C1-18 alkyl

alkyl (substituted with OH), C5-8 cycloalkyl]. Thus, a composition comprising block

k
polypropylene 95, Youmex 1001 (maleated polypropylene) 5, Ca stearate
0.05, tetrakis[3-(3,5-di-tert-butyl-4hydroxyphenyl)propionyloxymethyl]methane 0.1,
tris(2,4-di-tert-butylphenyl) phosphite 0.1, and I 0.2 part was
injection-molded to give a test piece showing initial yellowness index
23.0 and cracking time 1104 h in a sunshine weathering test (83°, no raining). 705257-84-7

/USES /-84-/ RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (stabilizer; hindered alkoxyamine-containing acidic polymer compns.

with

RN

long-lasting weather resistance)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 19 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

ANSWER 20 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1371532 Document No. 151:338217 Superior light stabilization using a novel hindered amine light stabilizer. Negishi, Yoshinori, Kawamoto, Naoshi; Yukino, Toshinori (Polymer Additives R+D Laboratory, ADEKA Corporation, 5-2-13, Shirahata, Minami-ku, Saitama City, Saitama, 336-0022, Japan). Addcon World 2007, International Plastics Additives

Compounding Conference, 13th, Frankfurt, Germany, Sept. 5-6, 2007, 9/1-9/8. Smithers Rapra Technology Ltd.: Shrewsbury, UK. ISBN: 978-1-8475-5-018-3 (English) 2007. CODEM: 69KMT6.
The degradation of agricultural films made of polyethylene stabilized

with

hindered amine light stabilizer ADK Stab LA 81 and a conventional antioxidant package was investigated. The HALS showed excellent light stabilizing activity under acidic conditions of sulfur fumigation and sulfurous acid treatment.
705257-84-7, ADK Stab LA 81
RLi MOA (Modifier or additive use); USES (Uses)
(stabilization of polyethylene with hindered amine light stabilizers)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

L4 ANSWER 22 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2007:1173851 Document No. 147:450127 Rubber compositions with good
processability and their pneumatic tires. Negishi, Yoshinori; Ayabe,
Takashi; Tobita, Etsuo; Takeuchi, Takashi (Adeka Co., Ltd., Japan). Jpr
Kokai Tokkyo Koho JF 2007269911 A 20071018, 10pp. (Japanese). CODEN:
JKXXAF. APPLICATION: JF 2006-95351 20060330.

AB Title compns. contain 100 parts rubber and 0.01-20 parts
bis(1-oxy-2,2,6,6-tetramethylpiperidin-4-yl) carbonate (I). Thus,
4-hydroxy-1-oxy-2,2,6,6-tetramethylpiperidine was reacted with di-Me
carbonate in the presence of NaCMe to give I. A composition containing
100 parts

carbonate in the presence or NaLWE LO 92...

100 parts
natural rubber, 1 part I, 50 parts C black, 3 parts Zn flower, 2 parts
stearic acid, and 1 part antioxidant showed Mooney viscosity 58, 60, and
64 after kneading for 3 min, 5 min, and 7 min, resp.

IT 6146-58-3P
RL: IMF (Industrial manufacture); MOA (Modifier or additive use); PREP
(Preparation); USES (Uses)
(bis(oxytetramethylpiperidinyl) carbonate-containing rubber compns.

good processability for pneumatic tires)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

ANSWER 21 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN:
43139 Document No. 148:1446540 Process for the preparation of sterically hindered nitroxyl ethers. Schoening, Kai-Uwe; Fischer,

Walter:

er; Basbas, Abdel-Ilah; Dichtl, Alexander (Ciba Specialty Chemicals Holding Inc., Switz.). PCT Int. Appl. WO 2008003602 A1 20080110, 70 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR,

BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, FM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MK, MY, MZ, NN, NG, NI, NO, NZ, CM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SI, SM, SV, SY, TJ, TM, TN, TR, TT, TZ; RW: AT, BE, BF, JCF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, IT, LU, MC, ML, MR, MT, NE, NL, FT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2007-EP56292 20070625. PRIORITY: EP 2006-116619 20060705; EP 2007-102895 20070222.

The invention relates to a process for the preparation of a sterically energy and control of the corresponding sterically bindered mitroxyl

nitroxyl ether from the corresponding sterically hindered nitroxyl

nal by reacting it with a carbonyl compound and a hydroperoxide. The compds. prepared by this process are effective stabilizers for polymers against harmful effects of light, oxygen and/or heat, as flame-retardants for polymers and as polymerization regulators. Several nitroxyl radicals

polymers and as polymerization arguments.

alkylation with carbonyl compds. in the presence of a metal catalyst and hydroperoxide to give the corresponding nitroxyl ethers.

IT 705257-84-7P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of hindered nitroxyl ethers via radical alkylation of nitroxyl

radical with carbonyl compds. in the presence of hydroperoxide)
705257-84-7 CAPLUS
4-Piperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4\*-carbonate (CA INDEX NAME)

L4 ANSMER 23 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2007:1152194 Document No. 151:9581 Highly functional light stabilizer for
polyolefin. Ayabe, Takashi (Lab. for Resin Additive, ADEKA Co., Ltd.,
5-2-13 Shirohata, Minami-tu, Saitama-shi, 336-0022, Japan).
Purasuchikkusu, 58(9), 40-41 (Japanese) 2007. CODEN: PRSKAW. ISSN:
0555-7887. Publisher: Kopyo Chozakai.

AB The author has discussed on agricultural films which have high UV
stability, particularly polyolefin films, more particularly polyethylene
films, using specific highly functional light stabilizer such as ADK Stab
LA 81 manufactured by Asahi Denka. The paper has indicated that the
light

stabilizer (ADK Stab LA 81) has low basicity and therefore shows high light stability compared to conventional light stabilizers such as HALS. 705257-84-7
RL: MOA (Modifier or additive use); USES (Uses) (highly functional light stabilizer for polyolefin) 705257-84-7
CAPLUS 4-Piperidinol, 2, 2, 6, 6-tetramethyl-1-(undecyloxy)-, 4, 4'-carbonate (CA INDEX NAME)

ANSWER 24 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1941788 Document No. 147:2777280 Preparation of Phosphonium salts derivatives and their use in organic synthesis. Charette, Andre, Poupon, Jean-Christophe; Boezio, Alessandro (Valorisation-Rechereche, Can.). 2007:941788 Pat. Appl. Publ. US 20070197477 A1 20070823, 33pp., Cont.-in-part of

No. PCT/CA05/000523. (English). CODEN: USXXCO. APPLICATION: US 2006-539075 20061005. PRIORITY: US 2004-560592P 20040409; WO 2005-CA523 20050406

AB Title compds. are prepared and their use in organic synthesis is described.

Thus, reaction of (3-diphenylphosphinophenyl)triphenylphosphonium bromide (preparation given) with LiClO4 in MeCN gave 95% [(Ph2PC6H4)PPh3]+ClO4-

which

was used as reagent for Mitsunobu coupling reaction of menthol with

4-nitrobenzoic acid.

IT 867023-62-9P

RL: RCT (Reactant); SFN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)

(preparation of phosphonium salts derivs. and their use as reagent in

organic

synthesis)

synthesis)
867023-62-9 CAPLUS
1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[[[4(triphenylphosphonio)phenyl]methoxy]carbonyl]oxy]-, perchlorate (1:1)

CM 1

CRN 867023-61-8 CMF C35 H38 N O4 P

CM 2

CRN 14797-73-0 CMF C1 04

L4 ANSWER 24 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

867023-64-1P RL: RGT (Reagent); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation of phosphonium salts derivs. and their use as reagent in

(Continued)

organic

nic synthesis)
867023-64-1 CAPLUS
Phosphonium, [4-[[[[(1-hydroxy-2,2,6,6-tetramethyl-4-piperidinyl)oxy]arbonyl]oxy]methyl]phenyl]triphenyl-, perchlorate (1:1)
(CA INDEX NAME)

CRN 867023-63-0 CMF C35 H39 N O4 P

CM 2

CRN 14797-73-0 CMF C1 04

14 ANSWER 25 OF 67 CAPLUS COPYRIGHT 2009 ACS ON STN

2007:729700 Document No. 147:118641 Nitroxyl compound, and polymerization inhibitor and polymerization inhibitor composition using same. Negishi, Yoshinori; Tobita, Etsuo; Ayabe, Takashi (Adeka Corporation, Japan). PCT

Int. Appl. WO 2007074613 Al 20070705, 21pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BM, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, II, IN, IS, JP, RE, KG, RM, RN, KP, RR, RZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MM, MG, MK, MM, MW, MX, MZ, NA, NG, NI, NO, NZ, CM, FG, FH, FL, FT, FC, AB, GR, RT, BE, BF, BJ, CF, GG, GR, CI, CM, CT, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, TT, LU, MC, MI, MR, NE, NE, NT, FT, ST, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, TT, LU, MC, MI, MR, NE, NI, FT, SE, SN, TD, TG, TR. (Japanese). CODEN: PIXND2. APPLICATION: WO 2006—JP324377 20061206. PRIORITY: JP 2005—375673 20051227.

AB Disclosed is a compound having low volatility, high nitroxyl group concentration and excellent polymerization inhibitory activity. Also disclosed are a polymerization

polymerization
inhibitor and polymerization inhibitor composition using such a compound

disclosed are a nitroxyl compound represented by the formula (I) below, and

a polymerization inhibitor and polymerization inhibitor composition using such a nitroxyl

, agg. a nitroxy1 compound, e.g. bis(4-hydroxy-2,2,6,6-tetramethylpiperidinooxy) carbonate. 6146-58-31

RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses) ools (Uses) ..... (industrial manufacture); PREP (Preparation inhibitor composition using same)

RN 6146-58-3 CAPLUS
CN 1\_Directif ...

Gaing same)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 26 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2007;402380 Document No. 146:4225020 Piperidyl group-containing aren monomers, their (co)polymers with high NO content, and preparation thereof. Nanbu, Yoko; Taki, Takayuki (Adeka Co., Ltd., Japan). JF Kokai Tokkyo Koho JF 2007091811 A 20070412, 25pp. (Japanese). CC JKXXAF. AFFLICATION; JF 2005-280265 20050927. GT

 $^\star$  STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT  $^\star$ 

The title monomers are represented by I or II (R, R1 = O-liberating

AB The title monomers are represented by I or II (R, R1 = O-liberating group, carbamoyloxy), and are polymerized at -5-30° (in the presence of Ni complex catalysts) to afford the title polymers having unit III or IV (R2-R4 = O-liberating group, carbamoyloxy; m, k = 1-100; n = 0-100). polymers are useful for light stabilizers, conductivity-imparting agents, battery electrode materials, etc. Thus, 4-hydroxy-2,2,6,6-tetramethyl-1-piperidinyloxy was reacted with 5.11 g NaH in THF and then with 16.58 propargyl bromide in THF to give 4-propargyloxy-2,2,6,6-tetramethyl-1-piperidinyloxy, which was then reacted with 1-BuoK in DMSO at 35° to give 4-allyloxy-2,2,6,6-tetramethyl-1-piperidinyloxy (W) wis 90.0% yield. Then, V was polymerized in the presence of [(x-allyl)NiOCOCF3]2/PPh3 (prepared from dicyclopentadienylnickel and allyl triflate) at 25° to give a polymer (redox behavior given) with Mm 21,750 in 71.4% yield.

IT 934214-95-6P
RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation);

RL; IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT

(Reactant or reagent) (piperidyl-containing arene monomers forming nitrosyl-rich polymers useful

useful for light stabilizers or conductors)
RN 934214-95-6 CAPLUS
CN 1-Piperidinyloxy,
4-ethynyl-4-[(methoxycarbonyl)oxy]-2,2,6,6-tetramethyl(CA INDEX NAME)

ANSWER 27 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN:
191976 Document No. 144:2737550 Preparation of prodrugs containing novel biocleavable linkers. Satyam, Apparao (Nicholas Piramal India

India). U.S. Pat. Appl. Publ. US 20060046967 A1 20060302, 181 pp. (English). CODEN: USXXCO. APPLICATION: US 2005-213396 20050826. PRIORITY: US 2004-604632P 20040826; IN 2005-MU779 20050701. The invention provides compds. DI-L1-E-A-B-AI-E-(L-E-AI-B-A-E)0-2-L2-D2

is a bond, (CH2)1-6, (CH2CH2O)1-1000, S-S, S-S:O, S-SO2 or S-S:NH; A, Al are independently a bond, (CH2)1-8, 1,2-, 1,3- or 1,4-phenylene; D1 is a therapeutic agent having one or more functional groups OH, SH, NHR1,

CONHR1, OZCNHR1, SOZNHR1, SOZNHR1, NR1CONHNHR1 or NR1SOZNHR1 (R1 is H, alkyl, aryl, etc.); D2 is D1, a peptide, protein, monoclonal antibody, vitamin, NO, NO2, NONCate, a nitric oxide-releasing group, a polymer, etc.; E is independently CH2 or a bond, L1, L2 are independently a bond, O, S, NR1, L, or a linkage] or their pharmaceutically-acceptable salts

use as prodrugs, including NO-releasing prodrugs. Thus, aspirin prodrug 2-AcOC6H4CONHCH2CH2SSCH2CH2ONO2 was prepared and shown to release salicylate in rats in a sustained and controlled manner starting from 1 h through 12

h. 877865-31-1P RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THC (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(Uses)
(preparation of prodrugs containing novel biocleavable linkers)
877865-31-1 CAPLUS
1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[[[2-[[2(nitrooxy)ethyl]dithio]ethoxy]carbonyl]oxy]- (CA INDEX NAME)

ANSWER 28 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

CM 2

CRN 14797-73-0 CMF Cl 04

RL: RGT (Reagent); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of phosphonium salts derivs. and their use as solubility

CRN 867023-63-0 CMF C35 H39 N O4 P

CM 2

CRN 14797-73-0 CMF C1 04

ANSWER 28 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN: 1130651 Document No. 143:4060110 Phosphonium salts derivatives and their use as solubility controlling auxiliaries. Charette, Andre;

Foundation Control of the Control of

BZ, CA, CB, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, CD, GE, GB, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MM, MX, NA, NI, NA, NI, NO, NZ, CM, PG, PH, PL, FT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, IJ, TM, TN, TR, TI, TZ, UA, UG, US, CU, VN, YU, ZA, ZM, ZW, RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, RI, EL, IS, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: MO 2005-CAS23 2005406. PRIORITY: US 2004-5605529 200404095. The present invention relates to the use of compds. A3P or [A3P+L1]X- (A

various (un)substituted groups such as furyl, Ph, pyridyl, naphthyl, or
thiophenyl; X = anion; L1 = a linker, as solubility controlling
auxiliaries).
Thus, preparation of [3-Ph3P+C6H4FPh2]ClO4- is described in several steps
starting from 1,3-dibromobenzene, and was used as reagent for Mitsunobu
reaction of menthol with p-nitrobenzoic acid. These compds. can also be
used as solubility controlling fragments of a mol. The invention also

used as solubility controlling the solubility of a mol. or a substrate.

Moreover, the invention also relates to various phosphonium supported reagents or various phosphonium salts derivs.

IT 867023-62-9P
RL: RCT (Reactant); SFN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

INDEX NAME)

CM 1

CRN 867023-61-8 CMF C35 H38 N O4 P

L4 ANSWER 29 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2005;977018 Document No. 143:2872080 Weakly basic hindered amines having carbonate skeletons for synthetic zersin compositions and coating compositions with good long term stability and acid rain and chemical resistance. Negishi, Yoshinori; Ayabe, Takashi, Tobita, Etsuo (Asahi Denka Co., Ltd., Japan). PCT Int. Appl. WO 2005082852 A1 20050909, 39

DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW,

BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GB, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LK, LS, LT, LU, LV, MA, ND, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, CM, FG, PH, PH, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, RW, AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FT, FR, GA, GB, GR, IE, IS, TT, LU, MC, ML, MK, NK, NL, PT, SE, SN, TD, TG, TG, TGAPARSED. CODEN: PIXND2. APPLICATION: WO 2005-JP3807 20055028. PRIORITY: JP 2004-57297 20040302.

The present invention relates to hindered amines I, wherein R = C1-30 (hydroxy)alkpl or C2-30 alkenyl; n = 1-4 integer; R1 = C1-22 alkyl, C2-22 alkenyl, or II when n = 1; R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n =  $\frac{1}{2}$  R1 = C2-20 n-valent organic group when n = 0.00 n-valent org AB

Thus, 98.1 mmol 4-hydroxy-1-oxy-2,2,6,6-tetramethylpiperidine and 78.5 mmol dilauroyl peroxide were reacted for 6 h, 53.9 g 7.3% aqueous sodiv hydroxide solution and 25 g methanol was added into the resulting reaction

mixture to remove lauric acid, filtered, evaporated, 0.57 g sodium hydroborate

nydroporate
and reacted to remove 1-undecanoxy-2,2,6,6-tetramethylpiperidin-4-one to
give 4-hydroxy-1-undecanoxy-2,2,6,6-tetramethylpiperidine, 12.0 g of
which

n was reacted with 4.19 g diphenylcarbonate at 170-180° in the presence of potassium carbonate for 8 h to give bis(1-undecanoxy-2,2,6,6-tetramethylpiperidin-4-y1)carbonate, 0.5 parts

bis(1-undecanoxy-2,2,6,6-tetramethylpiperidin-4-yl)carbonate, 0.5 parts of which was mixed with PES 120 (LLDPE) 100, calcium stearate 0.05, tetrakis(3-(3,5-di-tetr-butyl-4-hydroxyphenyl)propionyloxymethyl)methane 0.05, and tris(2,4-d-terr-butylphenyl)phosphite 0.05 parts, kneaded, pelletized, and pressed to give a film, showing good acid rain resistance.

IT 705257-84-7P 863984-48-9P RL: IMF (Industrial manufacture); MOA (Modifier or additive use); PREP (Preparation); USES (Uses) (preparation of weakly basic hindered amines having carbonate skeletons for synthetic resin compns. and coating compns. with good long term stability and acid rain and chemical resistance)

RN 705257-84-7 CAPLUS

ANSWER 29 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued) 4-Fiperidinol, 2,2,6,6-tetramethyl-1-(undecyloxy)-, 4,4'-carbonate (CA INDEX NAME)

863984-48-9 CAPLUS
4-Piperidinol, 1-(heptadecyloxy)-2,2,6,6-tetramethyl-, carbonate (2:1)
(ester) (9C1) (CA INDEX NAME)

ANSWER 30 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

CM 1

CRN 861804-73-1 CMF C10 H18 N C

CM 2

CRN 9005-80-5

CMF Unspecified CCI PMS, MAN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

ANSWER 30 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1696785 Document No. 143:194667 Magnetic therapeutic nanoparticles comprising polymer-encapsulated superparamagnetic core and bioactive agents. Barry, Stephen E.; Sunderland, Christopher J.; Goodwin, Andrew (Alnis Biosciences, Inc., USA). PCT Int. Appl. WO 2005070471 A2 20050804 40 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LK, LS, LT, LU, LV, MA, MD, MG, MK, MI, MW, MC, MZ, NA, NI, NO, NZ, GM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, RN: AT, BE, BF, BJ, CF, GG, CB, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IS, IT, LU, MC, MI, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2005-US1755 20050120. PRIORITY: US 2004-537500P APPLICATION: WO 2005-US1755 20050120. PRIORIT: US 2004-05050.

20040120;
US 2004-616390P 20041006.

AB A magnetic therapeutic nanoparticle comprises a plurality of bioactive agents, a core of superparamagnetic material, and a polymeric encapsulant.

The magnetic therapeutic nanoparticles may be used for targeted bioactive agent delivery. A plurality of bioactive agent-containing magnetic therapeutic nanoparticles may be administered to the environment to be treated, and an encapsulated bioactive agent is released by application of sufficient amount of heat. Thus, iron pentacarbonyl (0.74 mL) was added a mixture of oleic acid (4.9 g) in octyl ether (28 mL) at 100° under N2, the solution was heated to 290° over 1 h, held at this temperature until it turned black, heated for further 30 min, and the mixture was cooled to room temperature Trimethylamine oxide (1.26 g) was added to the reaction mixture, the mixture was heated to  $130^{\circ}$  over 20 min, held at this temperature for h, heated to 290° over 1 h, and held at this temperature for 1 h. Washed and filtered product comprised oleic acid-stabilized maghemite nanoparticles (10 nm) which were exchanged with a tetraalkylammonium hydroxide, and reacted with ethylene oxide-methacrylic acid diblock copplymer, and cis-diamminedimitratoplatinum to obtain polymer-encapsulated magnetic therapeutic nanoparticles containing 4-6% platinum as a bioactive agent.

861804-74-2DP, reaction products with poly(ethylene glycol)
diacrylate-sodium acrylate copolymer
Ris SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological
study); PREP (Preparation); USES (Uses)
(nanoparticles coated with, magmetic therapeutic nanoparticles
comprising polymer-encapsulated superparamagnetic core and bioactive
agents)

861804-74-2 CAPLUS Inulin, 2,2,6,6-tetramethyl-1-oxy-4-piperidinyl carbonate (9CI) (CA INDEX NAME)

L4 ANSWER 31 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2001:712815 Document No. 135:2733550 Manufacture of 2,6-diethyl-2,3,6-trimethyl- and 2,2-diethyl-6,6-dimethyl-1-alkoxypiperidinyl carboxylate esters and carboxamides as radical polymerization initiators. Nesvadba, Peter; Zink, Marie-ddile; Kramer, Andreas (Ciba Specialty Chemicals Holding Inc., Switz.). Ger. Offen. DE 10113209 Al 20010927, 48 pp. (German). CODEN: GWXDEX. APPLICATION: DE 2001-10113209 20010319. PRIORITY: EP 2000-810246 20000322.

2005:696785

2,2-Diethyl-6,6-dimethyl-1-alkoxypiperidinyl carboxylate esters and carboxamides [I; R = COR1, O2CNHR1, NR2COR1, R2NCONHR1, OCR1NCOR3; R1 = AB Н,

CO2H, CO2(C1-4 alkyl), CO2Ph, C2-18 alkyl, C2-4 alkenyl, etc.; R2, R3 =

Н,

C1-18 alkyl; R1R3 with an N atom can form (un)saturated, (benzo-fused) 5-membered ring; R4 = CH2Ph, MeCHPh, Me2CPh, etc., with a proviso] are used as radical polymerization initiators for producing polymers with

used as radical polymerization initiators for producing polymers with narrow mol.

weight distribution. Seven title compds. were prepared and approx. 221 claimed. For example, adding 1.57 of AcCl to a solution of 6.36 g 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)piperidin-1-ol and 2.02 g Et3N in 50 mL PhMe at 0-5° and stirring the mixture for 2 h at 20° gave 6.9 g of a title ester II. Dissolving 338 mg II in 8 g Bu acrylate and heating for 5 h at 145° under Ar gave 6.64 g Bu acrylate oligomorer having number-average mol. weight 6700 and weight-average mol. weight 8700.

IT 362617-51-4 362618-68-6 362618-69-7 362618-53-6 362618-53-6 362618-53-1 RL: CAT (Catalyst use); USES (Uses) (manufacture of diethyltrimethyl- and diethyldimethyl(alkoxy)piperidinyl carboxylate esters and carboxamides as radical polymerization initiators)

RN 362617-51-4 CAPLUS

CN Carbonic acid, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4-piperidinyl methyl ester (CA INDEX NAME)

ANSWER 31 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued) L4

362617-52-5 CAPLUS
Carbonic acid, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4-piperidinyl ethyl ester (CA INDEX NAME)

362617-53-6 CAPLUS
Carbonic acid, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4-piperidinyl phenylmethyl ester (CA INDEX NAME)

RN

362617-54-7 CAPLUS Carbonic acid, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4-piperidinyl 1,1-dimethylethyl ester (CA INDEX NAME)

RN

362618-68-6 CAPLUS Carbonic acid, 2,2-diethyl-6,6-dimethyl-1-(1-phenylethoxy)-4-piperidinyl methyl ester (CA INDEX NAME)

L4 ANSWER 32 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
2001:564135 Document No. 135:1250350 Fire-resistant electrolyte solutions
and secondary nonaqueous electrolyte batteries. Yamada, Manabu; Kubota,
Nachiro (Denso Co., Ltd., Japan, Kashi Denka Kogyo K. K.). Jpn. Kokai
Tokkyo Koho JP 2001210365 A 20010803, 10 pp. (Japanese). CODEN:

APPLICATION: JP 2000-22245 20000131.

The electrolyte solns. contain an electrolyte salt dissolved in an

organic solvent, which contains a piperidine derivative I, where RO = C1-18 alkyl group; R1-4 = C1-4 alkyl groups; n = 1-6 integer; X = II or III; R = C2-10

0 alkenyl group, A = -0-, -NR5- or a single bond; R5 = CI-10 alkyl group; B = H or CI-10 alkyl group that may also have ether bonding, n-valent acyl group or carbamoyl group, -CO2(R6CCOO)mR7 (R6 = C2-6 alkylene group, R7 = CI-10 alkyl group that may also have ether bonding, or IV, m = 0 or I),

alkylene or oxydialkylene group connected to R5. The electrolyte solns. may also contain phosphate esters.
351331-30-1 351331-33-4
RL: MOA (Modifier or additive use); USES (Uses)
(fire resistant additives for electrolyte solns. in secondary lithium batteries)
351331-30-1 CAPLUS
Carbonic acid, ethyl 1-methoxy-2,2,6,6-tetramethyl-4-piperidinyl ester (CA INDEX NAME)

тт

L4 ANSWER 31 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

вM

362618-69-7 CAPLUS
Carbonic acid, 2,2-diethyl-6,6-dimethyl-1-(1-phenylethoxy)-4-piperidinyl
ethyl ester (CA INDEX NAME)

362618-70-0 CAPLUS Carbonic acid, 2,2-diethyl-6,6-dimethyl-1-(1-phenylethoxy)-4-piperidinyl phenylmethyl ester (CA INDEX NAME)

RN

362618-71-1 CAPLUS Carbonic acid, 2,2-diethyl-6,6-dimethyl-1-(1-phenylethoxy)-4-piperidinyl 1,1-dimethylethyl ester (CA INDEX NAME)

L4 ANSWER 32 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

351331-33-4 CAPLUS 4-Piperidinol, 1-methoxy-2,2,6,6-tetramethyl-, carbonate (2:1) (ester) (9CI) (CA INDEX NAME)

L4 ANSWER 33 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 2000:6600579 Document No. 133:1960040 Fire-resistant electrolyte solutions and secondary nonaqueous electrolyte batteries. Kubota, Naohiro; Takeuchi, Yasunori (Asahi Denka Kogyo K. K., Japan). Jpn. Kokai Tokkyo Koho JP 2000235867 A 20000829, 9 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1999-36258 19990215.

The electrolyte solms, contain electrolyte salts and organic solvents, which

includes N-oxy-2,2,6,6-tetramethyl-4-piperidine, preferably I [n = 1-6,

= trivalent C2-10 alkane radical, A = -O-, -NR2- or a single bond, R2 = C1-10 alkyl group, B = H or C1-10 alkyl group which may have ether bonding, n valent acyl or carbamoyl group, or -COO-(R3CCOO)mR4, R3 = C2-6 alkylene group, R4 = C1-10 alkyl group or II]. The electrolyte salt is selected from LiPF6, LiBF4., LiClO4, CF3SO3Li, (CF3SO2)ZNLi, (CF3SO2) 3CLi:

and the solvent may also contain III or IV (R5-8 = linear or branched C1 -4

(fluorinated) alkyl group, R9 = linear or branched C2-8 alkylene group, n = 0 or 1). 6146-58-3

IT 289499-43-0

6146-58-3 294994-43-U RI: DEV (Device component use); USES (Uses) (electrolyte solvent mixts. containing N-oxy-2,2,6,6-tetramethyl-4-piperidine derivs. and phosphorus compds. for secondary lithium batteries)

L4 ANSMER 34 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1996:640164 Document No. 126:46848 Original Reference No. 126:9229a,9232a Organic radicals exhibiting intermolecular ferromagnetic interactions

high probability: 4-arylmethyleneamino-2,2,6,6-tetramethylpiperidin-1-yloxyls and related compounds. Togashi, Kensuke; Imachi, Ron; Tomioka, Katsuyuki, Tsuboi, Hidenori; Ishida, Takayuki; Nogami, Takashi; Takeda, Naoya; Ishikawa, Masayasu (Dep. Appl. Phys. Chem., Univ. Electro-Communications, Chofu, 182, Japan). Bulletin of the Chemical Society of Japan, 69(10), 2821-2830 (English) 1996. CODEN: BCSJA8.

:
0009-2673. Publisher: Nippon Kagakkai.
A series of 4-arylmethyleneamino-2,2,6,6-tetramethylpiperidin-1-yloxyls
(4-arylmethyleneamino-TEMPO) and related compds. were synthesized, and
their magnetic susceptibility were measured by a SQUID magnetometer in

temperature range of 1.8-100 K. Of 165 radicals investigated, 52 kinds of

radicals exhibited intermol. ferromagnetic interactions. These were confirmed by the increase of effective magnetic moments in

low-temperature regions. Pos. Weiss temps. ( $\theta$ ), ranging from +0.03 to +0.75 K, were found for these materials. Over 100 kinds of radicals exhibited antiferromagnetic interactions with  $\theta$  ranging from -0.01 to -24 K. The surprisingly high probability of finding organic radicals with intermol.

mol. ferromagnetic interaction may be understood by the characteristic mol. arrangements in the crystals. An oxygen atom of an ND radical site of a piperidin-1-yloxyl molety is apt to locate near methyl- and/or methylene-hydrogens of  $\beta$ -positions of the adjacent mols., and the resultant spin polarization gives rise to parallel spin alignments of nearest ND sites in the crystals. 4-(-Iodophenylmethyleneamino)-TEMPO exhibited a bulk ferromagnetic transition at 0.4 K. Six radicals exhibited metamagnetic transitions at magnetic fields lower than 200 Oe below 0.1 K. 6146-58-3P

6146-58-39
RI: FNU (Preparation, unclassified); PRP (Properties); PREP (Preparation)
(4-axylmethyleneamino-2,2,6,6-tetramethylpiperidin-1-yloxyls and related organic radicals having high probability of intermol. ferromagnetic interactions)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

тт

ANSWER 33 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 6146-58-3 CAPLUS (Continued)

RN

Index Name) 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

289499-43-0 CAPLUS 1-Piperidinyloxy, 4-[(ethoxycarbonyl)oxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 35 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1995;364840 Document No. 122:213312 Original Reference No.
122:38991a, 38994a
Spin catalysis of the radical recombination reaction. Buchachenko,
Anatoly L.; Ruban, Lyudmila V.; Step, Eugene N.; Turro, Nicholas J.
(Institute of Chemical Physics, Russian Academy of Sciences, Moscow,
117334, Russia). Chemical Physics Letters, 233(3), 315-18 (English)
1995.

1995.

CODEN: CHPLBC. ISSN: 0009-2614. Publisher: Elsevier.
Rate consts. of the recombination of alkyl radicals (cyanoisopropyl and sec-phenethyl radicals) with nitroxide biradicals in dioxane and Et benzene exceed rate consts. with 'parent' mono radicals by 104-50%. The difference in reactivity is attributed to spin catalysis of the recombination reaction which occurs in the encounter pair of an alkyl radical with one of the biradical termini by the second spin-carrying terminus. 6146-58-3

6146-58-3 RL: PEP (Physical, engineering or chemical process); PRP (Properties);

(Reactant); PROC (Process); RACT (Reactant or reagent)
(rate consts. of recombination of alkyl radicals with nitroxide biradicals)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl-(CA INDEX NAME)

ANSWER 36 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 18358 Document No. 118:83580 Criginal Reference No. 118:1703a,1706a Light stabilizers for ambient cured coatings. Behrens, Rudolf A.; Mar, Andrew (Ciba-Geigy Corp., USA). U.S. US 5124378 A 19920623, 17 pp. Cont.-in-part of U.S. Ser. No. 99,420, abandoned. (English). CODEN: USXXMM. APPLICATION: US 1988-259945 19881019. PRIORITY: US 1987-99420 19870921.

USAAMA. APPLICATION: US 1980-259946 1980(119. PRIORIT: US 1987-3942)

AB Derivs. of N-acyloxy or hydrocarbyloxy hindered piperidines of specified structure are light stabilizers for ambient-cured coatings, giving good durability and weather resistance. A tung oil-modified phenolic resin containing 5 phr bis(1-acetoxy-2,2,6,6-tetramethyl-4-piperidinyl) sebacate (I)

gave coatings on red cedarwood with 60° gloss retention after 8 mo outdoor exposure 46.7%; vs. 24.2 without I, and 39.0 with bis(2,2,6,6-tetramethyl-4-piperidinyl) sebacate in place of I.

II 137452-96-1

RL USES (Uses)

(light stabilizers, for coatings)

RN 137452-96-1 CAPLUS

Carbonic acid, 1-(benzoyloxy)-2,2,6,6-tetramethyl-4-piperidinyl butyl ester (CA INDEX NAME)

useful as stabilizers for polymers other than polyolefins. A PVC plate containing 1% bis(1-methoxy-2,2,6,6-tetramethylpiperidin-4-yl) isophthalate (I), had AE value 2.8 (ASTM D-1925-63T) after exposuring for 3014 h in a weatherometer, vs. 6.7 without I.

IT 137452-96-1 RL: USES (Uses) (stabilizers, for polymers other than polyolefins)
RN 137452-96-1 CAPLUS
CN Carbonic acid, 1-(benzoyloxy)-2,2,6,6-tetramethyl-4-piperidinyl butyl ester (CA INDEX NAME)

ANSWER 37 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :512808 Document No. 117:1128080 Original Reference No. 117:10695a, 19698a Polyolefin compositions stabilized with N-hydrocarbyl(carbonyl)oxy-substituted hindered amines. Galbo, James P.; Seltzer, Raymond; Ravichandran, Ramanathan; Patel, Ambelal R. (Ciba-Gelgy Copr., USA). U.S. US 5096950 A 19920317, 18 pp. Cont.-in-part of U.S. Ser. No. 259,946, abandoned. (English). CODEN: USXXAM. APPLICATION: US 1990-562783 19900806. PRIORITY: US 1988-259946 19881019.
The title stabilizers are especially useful in polyolefin films for mhouses

greenhouses

or swimming pool covers. Polypropylene containing a cinnamate etabilizer was

ilizer was mixed with 0.1% di(1-butylcarbomoyloxy-2,2,6,6-tetramethylpiperidin-4-yl)2,2-diethylmalonate (I), was extruded to a film, and exposed to light to give time to failure 890 h, vs. 340 without I. 137452-96-1 RL: USES (Uses) (stabhlizers, to heat and light and oxygen, for polyolefins) 137452-96-1 CAPLUS Carbonic acid, 1-(benzoyloxy)-2,2,6,6-tetramethyl-4-piperidinyl butyl ester (CA INDEX NAME)

L4 ANSWER 39 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1991:82730 Document No. 114:827300 Original Reference No. 114:14149a,14152a
Peroxides containing hindered amine moieties with low basicity as
polymerization initiators. Seltzer, Raymond; Winter, Roland A. E.;
Schirmann, Peter J. (Ciba-Geigy A.-G., Switz.). Eur. Pat. Appl. EP
389423

389423

389423
Al 19900926, 19 pp. DESIGNATED STATES: R: DE, FR, GB, IT. (English). CODEN: EPXXDW. APPLICATION: EP 1990-810191 19900313. PRIORITY: US 1989-326353 19890321.
AB The title peroxides provide a polymer containing a hindered amine stabilizer chemical bonded to the polymer and the low basicity of the peroxides prevents interaction with acid catalysts used in some polymerization systems. Thus. Bu

, Bu acrylate 25, 2-hydroxyethyl acrylate 30, Bu methacrylate 27, styrene 15, and acrylic acid 3% were polymerized with 6.5 g (based on 100 g monomer

nre) OO-tert-amyl-O-(1-cyclohexyloxy-2,2,6,6-tetramethylpiperidine-4-yl) monoperoxycarbonate gave a polymer containing a hindered amine stabilizer (I)

Polypropylene containing 0.2% I was molded into test pieces exhibiting

failure resistance in a fluorescent sunlight/black light failure chamber. 132147-26-3 RL: USES (Uses)

(polymerization initiators, for vinyl polymers containing hindered amine

stabilizers)

Stabilizers)
32147-26-3 CAPLUS
Carbonoperoxoic acid, O-[1-(cyclohexyloxy)-2,2,6,6-tetramethyl-4-piperidinyl] OO-(1,1-dimethylpropyl) ester (9CI) (CA INDEX NAME)

14 ANSWER 40 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1983:575125 Document No. 99:175125 Original Reference No. 99:26849a,26852a
Use of the ESR half-field transition to determine the interspin distance
and the orientation of the interspin vector in systems with two unpaired
electrons. Eaton, Sandra S.; More, Kundalika M.; Sawant, Bhimrao M.;
Eaton, Gareth R. (Dep. Chem., Univ. Colorado, Denver, CO, 80202, USA).
Journal of the American Chemical Society, 105(22), 6560-7 (English) 1983.
CODEN: JACSAT. ISSN: 0002-7863.
AB For systems containing 2 unpaired electrons with g values .apprx.2 and an
interspin distance of r, the intensity of the forbidden half-field
transition is proportional to r-6. The hyperfine splitting of the
half-field signal depends on the relative orientations of the nuclear
hyperfine tensors for the 2 electrons and the orientation of the
interspin
vector. Thus, the r value and the relative orientations of the hyperfine
tensors were determined independently, and also independently of the
value of
the exchange coupling constant J. The method was calibrated with 7 mols.
with well-characterized geometries: a Cu dimer, 4 dinitroxyl radicale and
2 spin-labeled Cu complexes. The absolute value of J for these mols.

2 spin-labeled Cu complexes. The absolute value of J for these mols. ranged from .apprx.100 to 25 + 10-4 cm-1. Two 3-spin systems were examined The method was also applied to a spin-labeled Cu complex for which the Cu-nitroxyl distance could not have been obtained by other ESR methods. If 6146-58-3 RL: PRP (Properties)

(ESR half-field transition of, interspin vector in relation to) RN 6146-58-3 CAPLUS
CN 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 41 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

L4 ANSWER 41 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1982;519348 Document No. 97:119348 Original Reference No. 97:19657a,19660a Metal-nitroxyl interactions. 23. Dinitroxyl adducts of paramagnetic metal complexes. Sawant, Bhimrao M.; Eaton, Gareth R.; Eaton, Sandra S. (Dep. Chem., Univ. Denver, Denver, CO, 80208, USA). Journal of Magnetic Resonance (1969-1992), 45(1), 162-9 (English) 1981. CODEN: JCMRA4. ISSN: TSSN:

0022-2364.

0022-2364.

AB The characteristic EPR spectrum of a nitroxyl biradical with intermediate electron-electron exchange, changed to a 3-line mononitroxyl EPR spectrum upon coordination of anitroxyl O to the spin-1/2 metal complexes Cu(tfac)2, Cu(hfac)2, VO(tfac)2, or VO(hfac)2 (tfac = trifluoroacetylacetone, hfac = hexafluoroacetylacetonate). Similar behavior was observed for a nitroxyl biradical with large electron-electron exchange when a nitroxyl O was coordinated to VO(tfac)2 or VO(hfac)2.

One

One
electron from the dinitroxyl couples with the metal unpaired electron, yielding a net spin of 1/2. Thus the dinitroxyl-paramagnetic metal complex becomes a mononitroxyl. Approx. equilibrium consts. were obtained for these Lewis acid-Lewis base interactions.

IT 6146-58-3D, copper and vanadium complexes R1: PRP (Properties)
(ESR and equilibrium consts. for)
RN 6146-58-3 CAPLUS
CN 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

6146-58-3 IT

RE: PRP (Properties)

(ESR study of interaction of, with metal complexes)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA

INDEX NAME)

L4 ANSWER 42 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1982:21272 Document No. 96:21272 Original Reference No. 96:3551a,3554a Effect of electron spin exchange on the interaction of stable nitroxyl radicals with triplet states of cyanine dyes. Borisevich, Yu. E.; Kuz "min, V. A.; Kokorin, A. I.; Sennikov, G. P.; Novochlova, G. A.; Shapiro, A. B. (Inst. Khim. Fiz., Moscow, USSR). Izvestiya Akademii Nauk SSSR, Seriya Khimicheskaya (9), 2019-23 (Russian) 1981. CODEN: IASKA6. ISSN: 0002-3353.

(Continued)

The efficiency of quenching of the triplet state of a cyanine dye by stable nitroxyl biradicals, as studied by flash photolysis in EtOH at 20°, increased with increasing spin exchange between the paramagnetic centers of the biradical, accompanied by a change in the quenching mechanism (formation of a charge-transfer complex). The quenching rate constant increased from .apprx.1.3 + 107 to .apprx.3.0 + 107 L/mol-s as the absolute values of the spin-exchange integral measured in hyperfine interaction constant units increased from 0.2 to

IT

(quenching by, of cyanine dye triplet state)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA
INDEX NAME)

#### L4 ANSWER 42 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

ANSWER 43 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :192981 Document No. 92:192981 Original Reference No. 92:31201a,31204a The study of nitroxide radical active esters as spin labels on muscle protein actin. Belagyl, J.; Grof, F.; Pallai, G.; Tigyi, J. (Cent. Lab., Med. Univ., Pecs, H-7624, Hung.). Acta Biochimica et Biophysica emisae Scientiarum Hungaricae, 14(3), 183-8 (English) 1979. CODEN: ABBPAP. ISSN: 0001-5253.

Nitroxide radical active esters (I, II, III, and IV) were used to label muscle actin and to study the orientation dependence of the ESR spectra  $\alpha$ AB

the labeled protein. The labels were located at  $\geq 2$  different sites of the protein with strong polar environment and different mobilities. The ESR spectrum of the strongly immobilized labels exhibited orientation dependence, the N-O bond axis of the spin labels being nearly perpendicular to the long axis of the F-actin threads. The labels underwent a rapid rotational motion about an axis directed perpendicular to the filament axis. The application of nitroxide radical active esters may be useful in the study of ordered systems. 71645-08-4D, actin derivs.

RL: PRP (Properties)
(ESR of)
71645-08-4 CAPLUS
1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[[(4-nitrophenoxy)carbonyl]oxy]-

1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[[(4-nitrophenoxy)carbonyl]oxy]-(CA INDEX NAME)

L4 ANSWER 43 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

L4 ANSWER 44 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1979:558069 Document No. 91:1580690 Original Reference No. 91:25525a,25528a Nitroxyls; IV. Synthesis of spin-labeled N-(4-piperidinyloxycarbonyl)imidazoles and 4-piperidinyloxycarbonyl azides and their reaction with amino acid derivatives. Hankovszky, H. O.;

and their reaction with amelio accelerations. Hideg,
K.; Lex, L.; Tigyi, J. (Biophys. Dep., Univ. Pecs, Pecs, 7643, Hung.).
Synthesis (7), 530-1 (English) 1979. CODEN: SYNTBF. ISSN: 0039-7881.
OTHER SOURCES: CASREACT 91:158069.

AB Treating piperidin-1-oyl I (R = H) with carbonyldiimidazole in Et20-THF

at room temperature gave 76% imidazole II which was treated with p-Mec6H4SO3H in acetone to give 90% imidazolium tosylate whose treatment with NaN3 in H2O at room temperature gave 92% azide I (R = CON3) (III). III was also prepared in

66% yield by treating I (R = COC6H4NO2-p) with NaN3 in acetone/H2O at

room temperature Treating III with H-X-OR1 (X = Gly, R1 = Et; X = Phe, Trp, R1 = H)

H)

gave 65-80% I (R = (O-X-OR1).

71645-08-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with sodium azide)

71645-08-4 CAPLUS
1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[(4-nitrophenoxy)carbonyl]oxy](CA INDEX NAME)

ANSWER 45 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 189431 Document No. 88:89431 Original Reference No. 88:14011a,14014a Synthesis of nitroxyl derivatives of benzo[a]pyrene. Schlude, H. (Abt. Org. Chem. Spektrosk., Max-Planck-Inst. Biochem., Martinsried, Fed. Rep Ger.). Organic Preparations and Procedures International, 9(6), 289-96 (English) 1977. CODEN: OPPIAK. ISSN: 0030-4948.

Me.

R1(CH2)nOCO2R (R = benzo[a]pyren-6-yl, R1 = I, n = 0 or 1 throughout this abstract) were prepared by treating R1(CH2)nOH with COCl2, then treating

obtained chloroformate with ROAc. Also prepared were R1CO2R,

obtained chloroformate with ROAc. Also prepared were RICO2R,
RCO2 (CH2)RR1,
RCH (CH)R1, RCO2CO2Et, and RCOR2 (R2 = 1H-imidazol-1-y1).

7 65693-98-3P
RLI SPN (Synthetic preparation); PREP (Preparation)
(preparation and condensation with benzopyrenyl acetate)
RN 65693-98-3 CAPLUS
CN 1-Piperidinyloxy, 2,2,6,6-tetramethyl-4-[[[(2,2,6,6-tetramethyl-1-oxy-4-piperidinyl)carbonyl]oxy]carbonyl]oxy]- (9CI) (CA INDEX NAME)

IT

65694-06-6P
RL: SFN (Synthetic preparation); PREP (Preparation)
(preparation of)
65694-06-6 CAPLUS
1-Piperidinyloxy, 4-[[(benzo[a]pyren-6-yloxy)carbonyl]oxy]-2,2,6,6tetramethyl- (9CI) (CA INDEX NAME)

L4 ANSWER 46 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1977.424186 Document No. 87:24186 Original Reference No. 87:3847a,3850a
Acid esters of 4-piperidinol derivatives and their use as stabilizers.
Murayama, Keisuke; Morimura, Shoji; Yoshioka, Takao; Horiuchi, Hideo;
Higashida, Susumu (Sankyo Co., Ltd., Japan). Can. CA 997353 19760921, 32
pp. (English). CODEN: CAXXA4. APPLICATION: CA 1975-237404 19751010.
AB Methylated aza[5.5]spiroundecanes and acid esters of 4-piperidinol as

prepared in the presence of alcoholysis catalyst and used as heat—and light—stabilizers for polyolefins, PVC [9002-86-2], polyamides, and polyurethanes. Thus, 4-hydroxy-2,2,6,6-tetramethylpiperidine [2403-88-5] was heated with Bz0ft [93-89-0] in xylene containing NaOH to give 4-benzoyloxy-2,2,6,6-tetramethylpiperidine [26275-88-7], and tetrakis (2,2,6,6-tetramethyl-apperidyl) pyromelitate [39111-20-1] was prepared similarly from the corresponding secondary alc. and added (0.25 parts/100 parts resin) to CM 1011 (nylon 6) [2503-84-4]. A film from this composition exhibited elongation and tensile strength retentions 73

IT

and

70%, resp., after a 300-h exposure to UV at 45°, and 73 and 76%, resp., after a 2-h aging at 160°.
6146-58-3
RL: USES (Uses)
(heat- and light-stabilizers, for plastics)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

ANSWER 45 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

L4 ANSWER 47 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1977;121125 Document No. 86:121125 Original Reference No. 86:19123a,19126a Crystal and molecular structure of a stable biradical of bis(2,2,6,6-tetramethylpiperidin-1-oxyl) carbonate C19H34N2O5. Shibaeva, R. P.; Lobkovskaya, R. M.; Rozenberg, L. P. (Otd. Inst. Khim. Fiz., Chernogolovka, USSR). Zhurnal Strukturnoi Khimii, 17(5), 876-80 (Russian)
1976. CODEN: ZSTKAI. ISSN: 0136-7463.

X-ray anal. of I yielded crystallog. data, bond lengths, and bond angles. The piperidine rings had the chair conformation. The N-O bond (1.287 Å) made an angle of 19.4° with the CNC plane. The average distance between the paramagnetic centers was 11.58 Å, whereas the intermol. distance was only 6.0-6.5 Å. GH46-58-3 RL: PROC (Process) (x-ray anal. of)

IT

(x-ray anal. of) 6146-58-3 CAPLUS

1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

ANSWER 48 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
:42840 Document No. 82:42840 Original Reference No. 82:6809a,6812a
Determination of the distance between the paramagnetic fragments in
biradicals from the forbidden transition AMs = 2. Dubinskii, A. A.;
Grinberg, O. Ya.; Tabachnik, A. A.; Shapiro, A. B.; Ivanov, V. P.;
Rosantsev, E. G.; Lebedev, Ya. S. (Inst. chem. Phys., Moscow, USSR).
Biofirika, 19(5), 840-2 (Russian) 1974. CODEN: BIOFAI. ISSN: 0006-3029.
A method for calcn. of radical-radical distances in diradicals was
developed, based on the relative intensities of the EPR spectra of the
forbidden transition AMs = 2; calcns. were given for 7 piperidinooxy
diradicals.
6146-58-3
RL: PRP (Properties)
(radical-radical distance in, calcn. for)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA
INDEX NAME)

тт

14 ANSWER 50 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1974:424880 Document No. 81:24880 Original Reference No. 81:4021a,4024a
Intramolecular exchange and dipole-dipole interactions in solutions of some iminoxyl biradicals. Kokorin, A. I.; Parmon, V. N.; Suskina, V. I.;
Ivanov, Yu. A.; Rozantsev, E. G.; Zamaraev, K. I. (Inst. Khim. Fiz.,
Moscow, USSR). Zhurnal Fizicheskoi Khimii, 48(4), 953-6 (Russian) 1974.
CODEN: ZFKHA9. ISSN: 0044-4537.

AB Dipole-dipole interactions and the exchange integrals were calculated from the

AB Dipute-upon .....

From the EPR spectra of diesters of 4-hydroxy-2,2,6,6-tetramethylpiperidinoxy

H2SO2, H2SO3, H2SO4, H2CO3, and RP(O)(OH)2 (R = Ph, vinyl, styryl, and  $\beta$ -chlorostyryl), and correlated with the distances between N-O-groups. The exchange integrals depended more on the nature of the ester central atom than on the nature of the substituents on it. 6146-58-3 RL: PRP (Properties) (dipole-dipole interactions and intermol. exchange in, EPR in relation

to) 6146-58-3 CAPLUS

1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

ANSWER 49 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :551152 Document No. 81:151152 Original Reference No. 81:23565a,23568a Spin exchange in nitroxyl biradicals. Metzner, E. Kurt; Libertini, Louis J.; Calvin, M. (lawence Berkeley lab., Univ. California, Berkeley, CA, USA). Journal of the American Chemical Society, 96(20), 6515-16 lish)

lish)
1974. CODEN: JACSAT. ISSN: 0002-7863.
For diagram(s), see printed CA Issue.
The electron spin exchange energy was measured as a function of solvent and of temperature for 3 nitroxyl biradicals (I,II,III). The effect of

and of temperature for 3 nitroxyl biradicals (I,II,III). The effect of errature
can be explained in terms of the flexibility of the mols. The dependence of the exchange on solvent is complex and not readily interpretable; however, it seems to be related to solvent polarity and the structure of the bridge between radical subunits.
6146-58-3
RL: PRF (Properties)
(spin exchange energy in, solvent and temperature effects on)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

14 ANSWER 51 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1974:121793 Document No. 80:121793 Original Reference No. 80:19616h,19617a
Polyolefins containing 4-piperidinol esters as uv stabilizers. Murayama,
Keisuke; Morimura, Shoji; Yoshioka, Takao; Horiuchi, Hideo; Higashida,
Susumu (Sankyo Co., Ltd.). Jpn. Kokai Tokkyo Koho JP 48072240 19730929
Showa, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1972-42959
19720428.

AB Polymer stabilizers were 4-piperidinol esters I [R, R1 = alkyl, RR1 =
saturated alicyclic member, CH2CMe2NR2CMe2CH2, R2 = 0 radical, lower
alkyl, R3
= aliphatic or aromatic acyl (n=1), diacyl (n=2), triacyl or trivalent
P(O) or P
(n=3), tetraacyl or Si (n=4)]. For example, a high d. polyethylene
[9002-88-4] sheet (0.5 mm thickness) containing 0.25 phr
1-ara-4-benzoyloxy-1,2,2-trimethylspiro[5.5]undecane I [RR1 = (CH2)5, R2

Me, R3 = Bz, n = 1] [51249-12-8] had uv resistance (time to brittle in a fadeometer at  $45.\deg$ .) 1800 hr, compared with 400 hr for a film not

containing
the stabilizer.
IT 6146-58-3
RL: USES (Uses)

(stabilizers, toward uv light, for polyolefins)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 52 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

1974:107500 Document No. 80:107500 Original Reference No. 80:17283a,17286a
Evidence for slow exchange in ESR spectra of nitroxide biradicals.
Parmon, V. N.; Kokorin, A. I.; Zhidomirov, G. M.; Zamaraev, K. I. (Inst. Chem. Phys., Moscow, USSR). Molecular Physics, 26(6), 1565-9 (English)
1973. CODEN: MOPHAM. ISSN: 0026-8976.

AB The ESR spectra of birs(2,26,6-tetramethylpiperidinol-1-oxyl) sulfide (I)
and carbonate (II) were measured in PhMe solns. at 17-71°. The ESR
spectra of II were also measured in CC14, CG86, EtoB, Me2CO, and PhMe at
25°. The spectra of biradicals I and II were attributed to fast
and slow exchange, resp.

IT 6146-58-3

EL: PRP (Properties)
(ESR of, slow exchange in)
RN 6146-58-3 CAPLUS

CN 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA
INDEX NAME)

L4 ANSWER 54 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1972:78698 Document No. 76:78698 Original Reference No. 76:12651a,12654a Nitroxides. XLVI. Determination of the N-O stretching frequency in piperidinic nitroxide free radicals. Morat, C.; Rassat, A. (Lab. Chim. Crg. Phys., C.E.N. Grenoble, Grenoble, Fr.). Tetrahedron, 28(3), 735-40 (French) 1972. CODEN: TETRAB. ISSN: 0040-4020.
AB By a comparison of the ir spectra of piperidinic nitroxide radicals with spectra of the corresponding 15N-labeled radicals, the ir frequency of the

N-O stretching vibration was measured.
6146-58-3
RL: PRP (Properties)
(ir spectrum of)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethylINDEX NAME)
(CA

L4 ANSWER 53 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1972:565545 Document No. 77:165545 Original Reference No. 77:27191a,27194a 4-Piperidinol derivatives as stabilizers for polymers. Murayama, Keisuke:

uke, Morimura, Syoji; Yoshioka, Takao; Horiuchi, Hideo; Higashida, Susumu (Sankyo Co., Ltd.). Ger. Offen. DE 2204659 19720810, 35 pp. (German). CODEN: GWXXEX. APPLICATION: DE 1972-2204659 19720128. Twenty-five title compds. [1, n = 1, R = H or Me; R1 = R2 = Me; R1R2 = (GR2)5, CH2CMe2NHCMe2CH2, or CH2CMe2NMeCMe2CH2; R3 = Ac, n-C17H35CO, Bz, p-C1C6H4CO, p-H2NC6H4CO, or  $\beta$ -C10H7CO; or n = 2-4; R = H; R1 = R2 = Me; R3 = CO, COCO, CO (GH2)2CO, CO (CH2)4CO, o - and p-(CC)4C6H4, P, PO, 1,3,5-(CC)3C6H3, Si, or 1,2,4,5-(CC)4C6H2] or their N-oxides were ared

prepared

by reaction of I (n = 1, R3 = H) with R3(OR4)n (R4 = Me or Et). I were
used at 0.2-0.5% concns. in polymers, e.g. Geon 103 EP [poly(vinyl
chloride)] [9002-86-2] or polyethylene (II) [9002-88-4], as heat and

stabilizers. Thus, a small amount KOH was added to I (n = 1, RR2 = Me,

H) and BzOEt in xylene, the mixture heated at 120.deg. with complete

removal

of EtOH formed, and kept 2 hr at 140.deg. to give

4-benzoyloxy-1,2,2,6,6-pentamethylpiperidine (III) [16597-34-5]. Plates
(0.5 mm thick), made from II containing 0.25% III were heated at 45.deg.

under

uv irradiation The samples became brittle after 1420 hr as compared

with 400
hr for II containing no III.

IT 6146-58-3
R1: USES (Ises)
(light stabilizers, for plastics)
RN 6146-58-3 CAPLUS

CN 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 55 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1971:456467 Document No. 75:56467 Original Reference No. 75:8903a,8906a
Asymmetric line broadening in the electron resonance spectra of
biradicals. Luckhurst, G. R., Pedulli, G. F. (Dep. Chem., Univ.
Southampton, Southampton, UK). Molecular Physics, 20(6), 1043-55
(English) 1971. CODEN: MOPHAM. ISSN: 0026-93% ich dets. the linewidths
in the electron resonance spectra of flexible biradicals is modulation of
the scalar electron-electron exchange interaction. In systems of high
viscosity, the modulation of the exchange interaction is often quenched,
and the rotational modulation of the anisotropic magnetic interactions
now

now constitutes the principal relaxation mechanism. A theoretical expression is derived for the broadening which results from this relaxation process. The applications of the theory to the determination of mol. configurations, electron-electron separations, and the sign of the exchange interaction are illustrated by comparison with the electron resonance spectrum of bis(2,2,6,6-tetramethyl-piperidinol-1-oxyl)carbonate. The theory is also of value in understanding the spectra of partially immobilized biradical spin labels.

IT 6146-50-3
RL: PRP (Properties)
(electron spin resonance of, line-broadening in relation to structure of)

of) 6146-58-3 CAPLUS

OLHO-50-5 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA
INDEX NAME)

ANSWER 56 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :116329 Document No. 72:116329 Original Reference No. 72:20967a,20970a Symmetric linewidth variations in the electron resonance spectra of biradicals. Luckhurst, Geoffrey R.; Pedulli, G. F. (Dep. Chem., Univ. Southampton, Southampton, UK). Molecular Physics, 18(3), 425-8 (English) 1970. CODEN: MOFBAM. ISSN: 0026-8976. Linewidth variations exhibited by the nitroxide biradical bis(4-hydroxy-2,2,2-6,6-tetramethylpiperidinooxy) carbonate were reported. The singlet and triplet lineqidths increased with increasing erature, and although the singlet lines never sharpened, the triplet lines decreased

width at .apprx.410°K. The results, which indicated that the spectral d. must increase with increasing temperature, were interpreted by using
a 2-configuration model.

a 2-configuration model.
6146-58-3
RL: PRP (Properties)
RL: PRP (Properties)
(electron spin resonance of, symmetric line width variations in)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSMER 58 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1968:456158 Document No. 69:56158 Original Reference No. 69:10495a,10498a
Electron paramagnetic resonance spectroscopic study of nitroxide mono-

bi-radicals. Lemaire, Henri Commis. Energ. At. (Fr.), Rapp., CEA-R

The hyperfine E.P.R. spectrum of nitroxide monoradicals is dominated by magnetic interaction of electrons with the N nucleus. Only the magnitude of the isotropic, or contact, interaction is measurable due to free tumbling. This magnitude depends on the structure and is altered by the solvent. The anisotropic, or dipole, interaction gives the values of the hyperfine tensor and the g-factor. For nitroxide bi-radicals, the hyperfine spectrum also depends on an exchange between singlet and let triple

states of the dimer. The sign of this exchange can be evaluated. In both

mono- and biradicals, increased solvent viscosity causes line-broadening. 6146-58-3 RL: PRP (Properties)

(electron spin resonance of, exchange in relation to)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

L4 ANSWER 57 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1968:463449 Document No. 69:63449 Original Reference No. 69:11867a,11870a
Is the terephthalic acid diester of
4-hydroxy-2,2,6,6-tetramethyl-piperidinooxy a strong [spin] exchange or
weak [spin] exchange biradical. Lemaire, H.; Rassat, A.; Rey, P.;
Luckhurst, G. R. (Lab. Chim. Org. Phys, C.E.N., Grenoble, Fr.).
Molecular

Pular Physics, 14(5), 441-7 (French) 1968. CODEN: MOPHAM. ISSN: 0026-8976. For diagram(s), see printed CA Issue. E.P.R. spectra were measured at room temperature on 5+10-4M solns. of  $\bullet RR$  (i,),  $p-[\bullet RC$ (i0)]2C6H4 (II),  $\bullet RC$ (i0)CH:CH:C(i0)R $\bullet$  (III),  $\bullet RC$ (i0)(CH2)4C(i0)R $\bullet$  (IV), and  $\bullet RC$ (i0)CR $\bullet$  (IV) in Me2NCHO. II and III exhibited spectra with 3 equal lines; IV, 5 lines characteristic of strong spin exchange; V was intermediate. The ratios

electron resonance line intensities of I to those of II-V were 1.925, 2.202, 0.584, and 0.669 for II, III, IV, and V (0.666 is the theoretical ratio for intermediate or strong exchange). II and III exhibit weak spin exchange; IV, strong spin exchange: V, intermediate spin exchange. 6146-58-3
RL: PRP (Properties)
(electron spin resonance of) 6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

14 ANSWER 59 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1968:414757 Document No. 69:14757 Original Reference No. 69:2791a
Nitroxides. XXVI. Effect of temperature on the spectral hyperfine
structure of the nitroxide biradicals. Lemaire, Henri; Rassat, Andre;
Rey, Paul (Lab. Chim. Org. Phys., C.E.N., Grenoble, Fr.). Bulletin de la
Societe Chimique de France (3), 886-92 (French) 1968. CODEN: BSCFAS.
ISSN: 0037-8968.

AB The biradicals for study were prepared by treating 2 mols.
2,2,6,6-tetramethyl-4-piperidinol 1-oxide with 1 mol. of a diacyl halide.
The esters prepared were carbonate (I), fumarate (m.184°), oxalate
(II), succinate (III), glutarate (m. 107°), adipate, pimelate,
suberate (m. 126°), sebacate, terephthalate (IV), and
2-norbornene-trans-5,6-dicarboxylate (m. 192°). All except II were
purified by chromatog. in CGH6 on Al203 (activity III). E.P.R. spectra
were obtained for solns. in HCONMez. Temperature was controlled to
t2°. Throughout the range of - 100° to +130°, IV
showed but 3 lines separated by 15.6 oe., the same as for the parent
radical.

ral. Little or no exchange is evident. I on the other hand shows a hyperfine splitting, with 7 major lines and 6 satellites. All other mols. showed a variation with temperature As an example, III at low temperature (-56°) lines separated by 15.6 oe.; a 2nd set of lines begins to appear at

(electron spin resonance of, conformation in relation to)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA

INDEX NAME)

ANSWER 60 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN :486393 Document No. 67:86393 Original Reference No. 67:16266h,16267a Spin exchange in nitroxide biradicals. Glarum, Sivert H.; Marshall,

H. (Bell Telephone Labs., Inc., Murray Hill, NJ, USA). Journal of Chemical Physics, 47(4), 1374-8 (English) 1967. CODEN: JCPSA6. I:

Chemical Physics, 47(4), 1374-6 (English) 1967. CODEM: JCPSN6. ISSN: 0021-9606.

The dependence of the electron spin-exchange energy on temperature and

ent has been studied for the carbonate, oxalate, and succinate diesters of tetramethylpiperidinoloxy. An analysis is given of exchange effects or E.S.R. line positions, intensities, and widths. For the carbonate

rs
through an indirect process involving the core of o electrons,
whereas at high temps. a direct exchange process is more important.
6146-58-3
EL: PRP (Properties)
(electron spin resonance of, spin exchange energy and)
6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA
INDEX NAME)

L4 ANSWER 61 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN

(Continued)

ANSWER 61 OF 67 CAPLUS COPYRIGHT 2009 ACS ON STN :463572 Document No. 67:63572 Original Reference No. 67:11918h,11919a Nitroxides. XX. Electron paramagnetic resonance of a nitroxide biradical: determination of the exchange sign. Lemaire, Henri (C.E.N., Grenoble, Fr.) Journal de Chimie Physique, 64(3), 559-71 (French) 1967. CODEN: JCQAY.

Grenoble, Fr.). Journal de Chimie Physique, 64(3), 559-71 (French) 1967. CODEN: JCPQAY.

GI For diagram(s), see printed CA Issue.

AB cf. 67: 43656u. E.P.R. spectra of dilute solns. of bis(2,2,6,6-tetramethyl-4-hydroxypiperidine l-oxide) carbonate (I) in dimethylformamide and in p-azoxyanisole were studied as a function of temperature For this biradical the electron-electron exchange interaction and the electron-lectron exchange interaction and the electron-luclei hyperfine interactions are of the same order of magnitude. In the isotropic phase the spectrum shows 6 lines due to 15N, each line having an electronic or a nuclear degeneracy. The exchange interaction, J, is found to be temperature dependent, due to conformational changes of I. Different conformations can be assumed either by inversion of one or both piperidine rings, or through their rotation around C-O bonds. The results are consistent with the lower-energy conformation having both hexaat. rings in the chair form and being stretched so that the distance between the 2 N-O groups is maximum. In the nematic mesophase of p-azoxyanisole where the electron-electron dipolar interaction is not averaged out, the degeneracy of the electronic transitions Ms = 0 0 Ms = 11 is removed, and the main lines of the isotropic spectrum are split into doublets. The results are discussed in terms of the anisotropic tensors of nitroxide monoradicals and of the conformation of I. The shift of the lines is used to determine the relative sign (opposite) of the singlet-triplet splitting J and of Dzz in the electron-electron dipolar tensor. Marked differences in the linewidths of various hyperfine lines result from temperature modulation, either of J or of the

hyperfine

lines result from temperature modulation, either of J or of the

anisotropies These differences provide a check on the sign of J and on the

conformation

of I as determined from the nematic phase results. The ground state of

biradical I is a singlet.

6146-58-3 RL: PRP (Properties)

(electron spin resonance of)
6146-58-3 CAPLUS
1-Fiperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethylINDEX NAME)

L4 ANSWER 62 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 1967:459414 Document No. 67:59414 Original Reference No. 67:11163a,11166a Temperature influence on the electron paramagnetic resonance spectrum o

nitroxide biradical. Lemaire, Henri; Levy, Bernard; Rassat, Andre (C.E.N., Grenoble, Fr.). Colloques Internationaux du Centre National de la Recherche Scientifique, Volume Date 1966, 164, 401-18 (French) 1967. CODEN: COINAV. ISSN: 0366-7634.

The E.S.R. spectrum of a nitroxide biradical, in which the J.vector.S1.vector.S2 exchange interaction is of the same order of magnitude as the A.vector.I1.vector.S1 isotropic interaction, is speed

composed of 6 lines when the N nucleus is 15N. The influence of temperature on  $\ddot{}$ 

spectrum of such a compound dissolved in HCONNe2 was studied. The splittings and the linewidths depend strongly on the temperature This is because the conformation of the biradical is temperature-dependent and

anisotropies of the Lande factor and the dipolar electron-nucleus and electron-electron interactions contribute to the linewidths. A crude theory suggests that a crossed term between the electron-electron dipole interaction and the anisotropy of the g factor explains the difference observed in linewidth between the  $Sz=0 \Leftrightarrow Sz=+1$  and  $Sz=0 \Leftrightarrow Sx=1$  transitions. 26 references.

RL: PRP (Properties)

(electron spin resonance of, hyperfine structure in, temperature and) 6146-58-3 CAPLUS 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA

INDEX NAME)

ANSWER 63 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 12881 Document No. 66:2881 Original Reference No. 66:667a,670a Inhibition of radical polymerization with nitroxide mono- and biradicals. Ruban, L. V.; Buchachenko, A. L.; Neiman, M. B.; Kokhanov, Yu. V. Vysokomolekulyarnye Soedineniya, 8(9), 1642-6 (Russian) 1966. CODEN: WMSDA8. ISSN: 0042-9368.

Exptl. behavior of the title radicals upon reaction with alkyl radicals

chain propagation are described. The structural formulas of both nitroxide mono- and biradicals are given. Kinetic measurements of the inhibiting effect of the monoradicals on styrene polymerization at 50°, initiated by acodisobutyronitrile, were carried out. The linear termination constant was found to be 2.1-3.2 + 104 1./mole/sec. (15%). In general, it was found that nitroxide monoradicals with different substituents have almost equal inhibiting effects. As for biradicals, a scheme for the recombination of alkyl radicals (like cyanoisopropyl and methylbenzyl) with nitroxide biradicals is suggested. The k2/k1 values for these biradicals are determined and tabulated. It

shown that the spin reactivity in the biradicals exceeds that in the monoradicals. This fact is explained on the basis of adiabaticity of the shown that the spin reactivity in the biradicals exceeds that in the monoradicals. This fact is explained on the basis of adiabaticity of trecombination. 6146-58-3 RL: USES (Uses) (Inhibitors, for polymerization of styrene) 6146-58-3 CAPLUS 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

14 ANSWER 65 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1966:35200 Document No. 64:35200 Original Reference No. 64:6455f-h,6456a-f
Nitroxides. XVII. Stable biradicals in the nitroxide series. Briere,
Roselyne; Dupeyre, Rose-Marie; Lemaire, Henri; Morat, Claude; Rassat,
Andre; Rey, Paul Bulletin de la Societe Chimique de France 3290-7
(French) 1965. CODEN: BSCFAS. ISSN: 0037-8968.

GI For diagram(s), see printed CA Issue.
Ab cf. preceding abstract Condensation of triacetonamine (I) with H2NNH2 in
refluxing (BCCH2CH2)20 gives 50% azine (II), m. 136°. Oxidation of II
with H2O2 in the presence of phosphotungstic acid gave a mixture
(separated by
chromatography on Al203) of the monoradical (V), m. 144°, and the
biradical (IV), m. 184°. With excess H2O2 in basic medium, the
yields are 23% IV and 47% V, for a 3-hr. stirring at ordinary
temperature IV
can also be obtained by the same oxidation process from V in aqueous
solution, or by

solution, or by
direct condensation of H2NNH2 to give III in refluxing (HOCH2CH2)20 in direct Condensation of Harmaz to give 111 in Perluxing (HACHZLHZ)20 II 16.6% yield. In the same way, condensation between 2 moles 2,22,5,5-tetramethyl-3-pyrrolidone (VI) and I mole H2NNHZ yields 43% corresponding azine (VII), m. 157°. Oxidation of VII by H2O2 in the presence of phosphotungstic acid gives, after 4 hrs., an orange precipitate, mixture of IX and X, separated by recrystn. in petroleum

where X only is soluble. The aqueous solution, ether-extracted, also gives

where X only is soluble. The aqueous solution, ether-extracted, also s crystals, mixts. of IX and X, separable by chromatography on Al2O3. IX, m. 198°, and X, m. 147°, are obtained in 40 and 19% yield, resp. IX can also be produced by condensation of 2 moles of the radical ketone VIII and 1 mole H2NNHZ in (HCCH2CH2)20 solution in 14% yield. Condensation of 2.2,6.6-tetramethyl-piperidin-4-ol 1-oxide (XI) with CCC12, (CCC1)2, or p-C1CO-C6H4CCC1 gives the corresponding carbonate

(XII) (m. 190°), oxalate (XIII) (m. 177°), or terephthalate (XIV) (m. 215°), obtained in 53%, 50%, and 69% yields, resp. IV and with the nitroxide 14N replaced by 15N (I = 1/2), have also be

starting from 15NH4Cl. With the biradicals prepared (IV, IX, XII-XIV), it

is possible to check the different predictions of theory, according to

relative magnitudes of the interaction J, between the 2 unpaired electrons, and the hyperfine splitting aN. If J « aN, the E.P.R. spectrum is composed of 3 equal lines, as in the monoradical (XIII, XIV). If J » aN, the hyperfine E.P.R. spectrum consists of 5 lines, with resp. intensities 1:12:32:1, as if each electron spent 1/2 the time on each N nucleus, the hyperfine splitting being reduced to aN/2: this is

case for IV, where the 7.40 oe. splitting is exactly 1/2 the hyperfine splitting measured for the monoradical V in the same solvent (Me2NCO), aN = 14.80 oe. The band widths (oe.) (2.40 for IV and 2.2 for V) are quite similar, which shows that dipolar interaction between the unpaired electrons contributes little to the band width in a solution with little viscosity. When the E.P.R. spectrum of IV with 15N is performed, the expected 1:2:1 triplet, due to interaction only with the nitroxidic N, is obtained with a splitting of 10.5 oe., normal for this isotopic substitution. The intermediate case which was also recorded shows the predicted outer satellites, 6 for 14N and 2 for 15N, in the E.P.R. tra

spectra of XII and its 15N derivative, with J/aN = 1.85; aN = 15.6 oe. (14N),

ANSWER 64 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN 160641 Document No. 64:160641 Original Reference No. 64:11401g-h,11402a Stabilization of polyurethans. (Toyo Spinning Co., Ltd.). NI. 6502388 19650827, 18 pp. (Unavailable). PRIORITY: JP 19640226. Polyurethans are stabilized against light and gases or vapors by mixing them (at any desired stage in the preparation or processing, but 1966:60641

them (at any desired staye in the first preferably in solution in a polar solvent) with 5-30 g./kg. of a phenylthiourea of

MIGHANHCSNRX', in which X is H, a halogen, or an alkyl, aryl, or aralkyl group; X' is H, alkyl, aryl or aralkyl; R is H or C1-4 alkyl; X and X'

contain halogen or C1-4 alkyl-substituted Ph; and (or) 0.3-20 g./kg. of

anhydride of a carboxylic acid with a dissociation constant at 25° of 10-4 to 10-5. Thus, a prepolymer was prepared by stirring 1 hr. at 85° of a mixture of 40 parts poly(tetramethylene ether) glycol (mol. weight 100) and 20 parts methylenebis(4-phenyl isocyanate). The product

dissolved in HCONMe2 (I) and cooled to 0°. A solution of 2 parts N2H4.H2O in 50 parts I was added, the mixture stirred for 15 min., and

reaction stopped by addition of 3 parts of a 10% solution of monochanolamine in

I. To 10 parts of the solution, 0.03 part diphenylthiourea was added,

the mixture extruded at room temperature into a dry spinning column at 180°. In a standardized burning test, the thread was rated 4 (5 meaning no discoloration, and 1 severe discoloration). The same sample, but containing no diphenylthiourea, was rated 1.
7392-64-5, Carbonic acid, phenyl ester, ester with 4-hydroxy-2,2,6,6-tetramethylpiperidinoxy

(polyoymethylene stabilization with polyamides and)
7392-64-5 CAPLUS
Piperidinooy, 4-hydroxy-2,2,6,6-tetramethyl-, phenyl carbonate (ester)
(8CI) (CA INDEX NAME)

ANSWER 65 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued) =  $1.32;~\rm AN = 21.90$  oe. (15N). The value of J depends upon both solvent (C6H6, MeCN, Me2SO, Me2NCHO, H2NCHO) and temp. 6146-58-3

(Derived from data in the 7th Collective Formula Index (1962-1966)) 6146-58-3 CAPLUS
1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

875835-02-2, Piperidinooxy, 4-hydroxy-2,2,6,6-tetramethyl-,

carbonate (ester)
(magnetic resonance absorption of)
875835-02-2 CAPLUS
Oxonium, [4-(carboxyoxy)-2,2,6,6-tetramethyl-1-piperidinyl]- (CA INDEX

L4 ANSWER 66 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN
1965:403240 Document No. 63:3240 Original Reference No. 63:574b-c New
stable iminoxyl radicals. Rozantsey, E. G.; Golubev, V. A.; Neiman, M.
B.; Kokhanov, Yu. V. (Inst. Chem. Phys., Moscow). Izvestiya Akademii

SSSR, Seriya Khimicheskaya (3), 572-3 (Russian) 1965. CODEN: IASKA6. ISSN: 0002-3353.

SSSR, Seriya Khimicheskaya (3), 572-3 (Russian) 1965. CODEN: IASKA6. ISSN: 0002-3353. 2,2,6,6-Tetramethyl-4-hydroxypiperidin-1-oxyl radical in C6H6 was treated with (CR2)c(NCO)2 and after heating 4 hrs. gave bis(2,2,6,6-tetramethyl-piperidin-1-oxyl) hexamethylenedicarbamate, pink, m. 115-16°, similarly, reactions with dichlorides of dicarboxylic acids in pyridine gave the corresponding esters of the above alc. with indicated acids: carbonic, m. 180°, oxalic, m. 179°, succinic, m. 141°, adipic, m. 180°, oxalic, m. 179°, succinic, m. 141°, adipic, m. 122°, pimelic, m. 90°, sebacic, m. 101°, terephthalic, m. 217°. Also reported was N,N'-bis(2,2,6,6-tetramethyl-4-piperidin-1-oxyl)-urea, m. 198-9°, prepared by oxidation of the dipiperidyl analog by means of MnO2. The E.P.R. spectra of the products were reported. The spectra showed changes in electronic interaction with changes in distances between the paramagnetic centers.
6146-58-3 CAPBUS
1-Piperiddinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)

875835-02-2P, Piperidinooxy, 4-hydroxy-2,2,6,6-tetramethyl-, carbonate (ester)
RL: PREP (Preparation)
(preparation of)
875835-02-2 CAPLUS
Oxonium, [4-(carboxyoxy)-2,2,6,6-tetramethyl-1-piperidinyl]- (CA INDEX

NAME)

L4 ANSWER 67 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN (Continued)

ANSWER 67 OF 67 CAPLUS COPYRIGHT 2009 ACS on STN:
403239 Document No. 63:3239 Original Reference No. 63:573f-h,574a-b
Condensation of 5-nitro-2-furaldehyde with 4-methyl- or 4-ethylpyridine.
Harada, Kinji; Emoto, Sakae (Inst. Phys. Chem. Res., Tokyo). Chemical &
Fharmaceutical Bulletin, 13(3), 389-91 (English) 1965. CODEN: CPBTAL.
ISSN: 0009-2363.
For diagram(s), see printed CA Issue.
The condensation of 5-nitro-2-furaldehyde (I) with 2-methyl- (II) and
4-methylpyridine (III) and with 2-ethyl- (IV) and 4-ethylpyridine (V) was
studied in order to obtain potential antibacterial agents. While
condensation of I with III and V on being heated 2 hrs. at 100° in
Ac2O has been reported (Belg. 615,319, CA 58, 11333h) to give VI and VII,
reinvestigation of these expts. has given only. apprx.108 yields crude VI
and VII, the purification of which was very difficult. Improved yields
were now obtained by using a mixture of AcOH and Ac2O as solvent. To
9.

4.7 g.

I and 3.2 g. III (dried 12 hrs. over NOH) in 20 ml. AcOH was added slowly during 20 mln. 5 ml. Ac2O at 60-70° with stirring, the solution heated 3 hrs. at 108° with stirring and cooled, the precipitate filtered off

filtrate (A) was kept] and extracted with 600 ml. hot MeOH, and the

evaporated in
vacuo, the residue poured into dilute HCl cooled in ice, and the
precipitate
filtered off and washed with saturated aqueous NaHCO3 and H2O gave crude

recrystn. of combined crude VI (3.5 g.) from MeOH gave 2.1 g. VI, m. 163-4°. To 2 g. I and 1.52 g. V in 8 ml. AcOH was added slowly 1.4 ml. Ac2C at 70° with stirring, the solution heated and stirred 3 hrs. at 110° and evaporated in vacuo, and the residue worked up like VI to give 1.18 g. VII, m. 150-1° (MeOH). Similar condensation of I with II and IV gave VIII, m. 176-7°, and IX, m. 199-9.5°, resp. To 1.5 g. I and 2 g. III (dried 12 hrs. over KOH) was added 6 ml. Ac2C (spontaneous temperature rise), the mixture let stand overnight at roomerature.

temperature,
heated 4 hrs. at 50-5°, and cooled, and the precipitate filtered off and
washed with 30 ml. Et20 to give 1.05 g. 5,5'-dinitro-2,2'-furoin

diacetate state (X), m.  $253-4^{\circ}$  (HCONMe2). This reaction also proceeded in the presence of V or 3-methylpyridine in lieu of III. VI-IX showed antibacterial action.

6146-58-3

(Derived from data in the 7th Collective Formula Index (1962-1966)) 6146-58-3 CAPLUS 1-Piperidinyloxy, 4,4'-[carbonylbis(oxy)]bis[2,2,6,6-tetramethyl- (CAINDEX NAME)

=>